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(54) Title: COMPOUNDS AND METHODS

(57) Abstract: This invention relates to substituted heterocyclic compounds which are modulators, agonists or antagonists, of the CCR5 receptor. In addition, this invention relates to the treatment and prevention of disease states mediated by CCR5, including, but not limited to, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, and inflammatory bowel disease, all in mammals, by the use of substituted heterocyclic compounds which are CCR5 receptor antagonists. Furthermore, since CD8+ T cells have been implicated in COPD, CCR5 may play a role in their recruitment and therefore antagonists to CCR5 could provide potential therapeutic in the treatment of COPD. Also, since CCR5 is a co-receptor for the entry of HIV into cells, selective receptor modulators may be useful in the treatment of HIV infection.

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COMPOUNDS AND METHODS

FIELD OF THE INVENTION

This invention relates to substituted heterocyclic compounds which are modulators, agonists or antagonists, of the CC chemokine receptor CC-CKR5 now designated as CCR5 (*Nature Medicine* **1996**, *2*, 1174-8). In addition, this invention relates to the treatment and prevention of disease states mediated by CCR5.

BACKGROUND OF THE INVENTION

T cells are not only key regulators of the immune response to infectious agents but are believed critical for the initiation and maintenance of the inflammatory reaction in a variety of chronic diseases. Increased numbers or enhanced activation state of T cells, especially CD4+ T cells, have been demonstrated in the synovium of individuals with rheumatoid arthritis (M.J. Elliott and R. N. Maini, Int. Arch. Allergy Immunol. 104: 112-1125, 1994), in the bronchial mucosa of asthmatics (C.J. Corrigan and A.B. Kay, Immunol. Today 13:501-506, 1992), in the lesions of multiple sclerosis (R. Martin and H. F. McFarland, Crit. Rev. Clin. Lab. Sci. 32: 121-182, 1995), in psoriatic lesions (J.L. Jones, J. Berth-Jone, A. Fletcher and P.E. Hutchinson, J. Pathol. 174: 77-82, 1994) and in the fatty streaks of atherosclerosis (R. Ross, Annu. Rev. Physiol. 57: 791-804, 1995).

T cells, as well as other inflammatory cells, will migrate into tissues in response to the production of a variety of chemotactic factors. Among these factors are a superfamily of 8-12 kDa proteins known as the chemokines. These proteins share structural features such as the presence of 3-4 conserved cysteine residues. RANTES, which stands for Regulated upon Activation Normal T cell Expressed and Secreted, is an 8 kDa protein member of CC branch of the chemokine family. These proteins recruit and activate immune and inflammatory cells through an interaction with G-protein coupled receptors. The CC branch is defined by the absence of an intervening amino acid residue between the first two cysteine residues and members of this family predominately elicit the migration of mononuclear cells, eosinophils and basophils (M. Baggiolini, B. Dewald, and B. Moser, Adv. Immunol. 55: 97-179, 1994; and J.J. Oppenheim, C.O.C. Zachariae, N. Mukaida, and K. Matsushima, Annu. Rev. Immunol. 9: 617-648, 1991).

RANTES potently produces chemotaxis of T cells, basophils, eosinophils, monocytes and mast cells. RANTES was originally identified as gene product induced late after antigen activation of T-cells (T.J. Schall, J. Jongstra, B.J. Dyer, J. Jorgensen,

et al., J. Immunol. 141:1018-1025, 1988), however, RANTES has been shown to be synthesized and secreted by a diverse group of cells that include epithelial and endothelial cells (C. Stellato, L.A. Beck, G.A. Gorgone, D. Proud, et al., J. Immunol. 155: 410-418, 1995; and A. Marfaing-Koka, O. Devergne, G. Gorgone, A. Portier, et al., J. Immunol. 154: 1870-1878, 1994), synovial fibroblasts (P. Rathanaswami, M. Hachicha, M. Sadick, T.J. Schall, et al., J. Biol. Chem. 268: 5834-5839, 1993) and dermal fibroblasts (M. Sticherling, M. Kupper, F. Koltrowitz, E. Bornscheuer, et al., (J. Invest. Dermatol. 105: 585-591, 1995), mesangial cells (G. Wolf, S. Aberle, F. Thaiss, et al., Kidney Int. 44: 795-804, 1994) and platelets (Y. Koameyoshi, A. Dorschner, A.I. Mallet, E. Christophers, et al., J. Exp. Med. 176: 587-592, 1992). In these cells, RANTES mRNA is rapidly upregulated in response to IL-1 or TNF•. Although RANTES mRNA is not usually detected in normal tissues (J.M. Pattison, P.J. Nelson, and A.M. Krensky, Clin. Immunother. 4: 1-8, 1995), increased mRNA or protein has been found in diseases characterized by a mononuclear infiltrate. For example, RANTES mRNA was visualized using *in situ* hybridization in renal allografts undergoing rejection (J.M. Pattison, P.J. Nelson, and A.M. Krensky, Clin. Immunother. 4: 1-8, 1995; and K.C. Nadeau, H. Azuma and N.I. Tilney, Proc. Natl. Acad. USA 92: 8729-8733, 1995) in the skin of atopic dermatitis patients after exposure to antigen (S. Ying, L. Taborda-Barata, Q. Meng, M. Humbert, et al., J. Exp. Med. 181: 2153-2159, 1995), and in endothelial cells of coronary arteries undergoing accelerated atherosclerosis after cardiac transplant (J.M. Pattison, P.J. Nelson, and A.M. Krensky, Clin. Immunother. 4: 1-8, 1995). Further, increased immunoreactive protein for RANTES has been detected in bronchoalveolar lavage fluid (R. Alam, J. York, M. Boyers, et al., Am. J. Resp. Crit. Care Med. 149: A951, 1994) and sputum from asthmatic individuals (C.M. Gelder, P.S. Thomas, D.H. Yates, I.M. Adcock, et al., Thorax 50: 1033-1037, 1995).

Several receptors have been identified that bind RANTES. In particular, CCR5, when expressed in either HEK 293 cells or CHO cells, binds RANTES. This receptor is expressed in T-cells and in monocytes and macrophages, immune/inflammatory cells which are important in the maintenance of a chronic inflammatory reaction. Pharmacological characterization of CCR5 indicates similarities to the RANTES binding site observed on isolated T cells. Therefore, antagonism of RANTES' action on CCR5, as well as antagonism of other natural modulators of CCR5, should inhibit the recruitment and activation of T cells and macrophages into inflammatory lesions and provide a novel therapeutic approach for the treatment of atopic and autoimmune disorders.

Since T cells express CCR5, selective receptor modulators of CCR5,

particularly antagonists, are likely to provide beneficial effects in diseases including, but not limited to, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, and inflammatory bowel disease, all in mammals, preferably humans. Furthermore, since CD8+ T cells have been implicated in chronic obstructive pulmonary disease (COPD), CCR5 may play a role in their recruitment and therefore antagonists to CCR5 could provide potential therapeutic activity in the treatment of COPD. Also, since CCR5 is a co-receptor for the entry of HIV into cells, selective receptor modulators may be useful in the treatment of HIV infection.

Compounds formula (I) having 5-$HT_{1D/1B}$ receptor antagonist activity have been reported in FR 2758328, published July 17, 1998; FR 2761069, published September 25, 1998; Matzen et al., *J. Med. Chem.* **2000**, *43*, 1149-1157; DE 197 56 036 A1, published June 24, 1999; WO 96/02525, published February 1, 1996; WO 97/28140, published August 7, 1997; WO 97/28141, published August 7, 1997; WO 98/31677, published July 23, 1998; U.S. Patent 5,789,412, issued August 4, 1998; WO 95/29907, published November 9, 1995; or compounds which inhibit leukotriene synthesis have been reported in WO 97/24328, published July 10, 1997; or compounds which antagonize tocolytic oxytocin receptor antagonist activity have been reported in WO 94/07496, published 14 April 1994, and WO 95/25443, published 28 September 1995.

Surprisingly, it has now been discovered that this class of non-peptide compounds, in particular substituted heterocyclic compounds of formula (I), function as CCR5 receptor modulators, and therefore, have utility in the treatment and prevention of disease states mediated by CCR5 receptor mechanisms.

SUMMARY OF THE INVENTION

The present invention is to compounds of formula (I), or a pharmaceutically acceptable salt, or solvate thereof, and their use as CCR5 modulators for the treatment and/or prophylaxis of certain disease states, including, but not limited to, COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, all in mammals, preferably humans. The preferred compounds for use as CCR5 modulators are those compounds of Formula (I) as noted herein.

In addition, the present invention is directed to a method of preventing or treating CCR5-mediated diseases in a mammal, preferably a human, by administering to the mammal an effective amount of a CCR5 receptor ligand, or a pharmaceutically acceptable salt or solvate thereof.

Further, the present invention is directed to methods for making and using the compounds of formula (I), as well as pharmaceutical compositions of formula (I) or a pharmaceutically acceptable salts or solvates thereof.

Yet further, the present invention is directed to the use of a CCR5 receptor ligand in the manufacture of a medicament for the prophylaxis or treatment of certain disease states, including, but not limited to, COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, for example in a mammal such as a human.

Still further, the present invention is directed to a CCR5 receptor ligand, or a pharmaceutically acceptable salt, or solvate thereof, for use in the prophylaxis or treatment of certain disease states, including, but not limited to, COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, for example in a mammal such as a human.

The present invention is also directed to combined therapy to prevent and treat inflammatory and immunoregulatory disorders or diseases, including asthma and allergic diseases, as well as rheumatoid arthritis and atherosclerosis, and those pathologies noted above, and is illustrated by the combination of the compounds of this invention and other compounds which are know for such utilities.

The present invention is further directed to combinations of the present compounds of formula (I) with one or more agents useful in the prevention or treatment of AIDS. For example, the compounds of this invention may be effectively administered, whether at periods of pre-exposure and/or post-exposure, in combination with effective amounts of the AIDS antivirals, immunomodulators, anti-infectives, or vaccines known to the skilled artisan.

DETAILED DESCRIPTION OF THE INVENTION

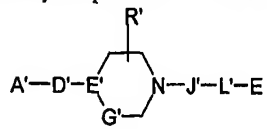
It has now been discovered that substituted heterocycles of formula (I) are CCR5 receptor modulators. It has also now been discovered that selective inhibition of

CCR5 receptor mechanisms by treatment with the receptor modulators of formula (I), or a pharmaceutically acceptable salt thereof, represents a novel therapeutic and preventative approach to the treatment of a variety of disease states, including, but not limited to, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, and inflammatory bowel disease, all in mammals, preferably humans. Furthermore, since CD8+ T cells have been implicated in COPD, CCR5 may play a role in their recruitment and therefore antagonists to CCR5 could provide potential therapeutic in the treatment of COPD. Also, since CCR5 is a co-receptor for entry into cells, selective receptor modulators may be useful in the treatment of HIV infection.

Compounds of formula (I) for use herein as CCR5 modulators include those compounds as described in FR 2758328, published 17 July 1998, FR 2761069, published 25 September 1998, WO 94/07496, published 14 April 1994, WO 95/25443, published 28 September 1995, and PCT/US00/01908, filed January 25, 2000. Each of these references is incorporated herein in their entirety.

Preferred compounds for use as CCR5 modulators are those compounds of formula (I) as noted herein.

A preferred group of compounds for use herein are those compounds of the formula (I) or a pharmaceutically acceptable salt or solvate thereof:



Formula (I)

in which:

the basic nitrogen in moiety E may be optionally quaternized with C_{1-6}alkyl or is optionally present as the N-oxide;

A' is aryl or heteroaryl, each of which is optionally substituted with one or more of $R^{1'}$; or A' is aryl or heteroaryl fused to a saturated or partly unsaturated 5-7-membered ring to form a higher order ring moiety, which ring moiety optionally contains 1 or 2 heteroatoms selected from oxygen, nitrogen or sulfur, wherein nitrogen may be optionally substituted with hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl; wherein the higher order ring moiety is optionally substituted with one or more of $R^{1'}$;

$R^{1'}$ is hydrogen, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, CH_2CF_3, aryl, aralkyl, $(CH_2)_a NR^{2'}R^{3'}$, $(CH_2)_a NR^{2'}COR^{4'}$,

$(CH_2)_a NR^{2'}CO_2R^{5'}$, $(CH_2)_a NR^{2'}SO_2R^{6'}$, $(CH_2)_a CONR^{7'}R^{8'}$, hydroxyC_{1-6}alkyl, C_{1-4}alkoxyalkyl (optionally substituted by a C_{1-4}alkoxy or hydroxy group), $(CH_2)_a CO_2C_{1-6}$alkyl, $(CH_2)_b OC(O)R^{9'}$, $CR^{10'}=NOR^{11'}$, $CNR^{10'}=NOR^{11'}$, $COR^{12'}$, $CONR^{7'}R^{8'}$, $CONR^{7'}(CH_2)_c OC_{1-4}$alkyl, $CONR^{7'}(CH_2)_a CO_2R^{13'}$, $CONHNR^{14'}R^{15'}$, $CONR^{7'}SO_2R^{16'}$, $CO_2R^{17'}$, cyano, trifluoromethyl, $NR^{2'}R^{3'}$, $NR^{2'}COR^{4'}$, $NR^{18'}CO(CH_2)_a NR^{18'}R^{19'}$, $NR^{18'}CONR^{18'}R^{19'}$, $NR^{2'}CO_2R^{5'}$, $NR^{2'}SO_2R^{6'}$, $N=CNR^{18'}NR^{18'}R^{19'}$, nitro, hydroxy, C_{1-6}alkoxy, OCF_3, hydroxyC_{1-6}alkoxy, C_{1-6}alkoxyC_{1-6}alkoxy, $OC(O)NR^{20'}R^{21'}$, $SR^{22'}$, $SOR^{23'}$, $SO_2R^{23'}$, $SO_2NR^{20'}R^{21'}$ or halogen, or $R^{1'}$ is a 5- to 7-membered ring containing 1 to 4 heteroatoms selected from nitrogen, oxygen, or sulfur, optionally substituted with hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, hydroxyC_{1-6}alkyl, (C_{1-6}alkyl)C_{1-6}alkyl, $CONR^{7'}R^{8'}$, $CO_2R^{17'}$, cyano, aryl, trifluoromethyl, nitro, hydroxy, C_{1-6}alkoxy, acyloxy, or halogen;

a' is 1, 2, 3 or 4;

b' is 0, 1, 2 or 3;

c' is 1, 2 or 3;

$R^{2'}$ and $R^{3'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{2'}$ and $R^{3'}$ together with the nitrogen to which they are attached, form a 5- to 6-membered heterocyclic ring which ring may be optionally substituted by an oxo group, or, when there are 6 ring members, the ring may optionally contain one oxygen or one sulfur atom;

$R^{4'}$ is hydrogen, C_{1-6}alkyl or C_{1-4}alkoxyalkyl, or, when $R^{1'}$ is $NR^{2'}COR^{4'}$, $R^{4'}$ is $(CH_2)_{1-3}$ and forms a ring with A';

$R^{5'}$ is C_{1-6}alkyl;

$R^{6'}$ is C_{1-6}alkyl or phenyl;

$R^{7'}$ and $R^{8'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{7'}$ and $R^{8'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring, wherein when there are 6 ring members, the ring may optionally contain one oxygen or one sulfur atom;

$R^{9'}$ is C_{1-4}alkyl, optionally substituted by a C_{1-6}alkoxy;

$R^{10'}$ and $R^{11'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{12'}$ is hydrogen or C_{1-6}alkyl;

$R^{13'}$ is hydrogen or C_{1-6}alkyl;

$R^{14'}$ and $R^{15'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{16'}$ is hydrogen or C_{1-6}alkyl;

$R^{17'}$ is hydrogen or C_{1-6}alkyl optionally substituted with one or more substituents selected from C_{1-6}alkyl, C_{1-6}alkoxy, hydroxy, or $NR^{2'}R^{3'}$;

$R^{18'}$ and $R^{19'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{20'}$ and $R^{21'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{20'}$ and $R^{21'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring which, when the ring is 6-membered, may optionally contain in the ring one oxygen or one sulfur atom.

$R^{22'}$ is hydrogen or C_{1-6}alkyl;

$R^{23'}$ is C_{1-6}alkyl;

D' is either a bond or represents $[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{a''}CO$, CO, SO_2, $CO[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}$, $S[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}OCO$, $NR^{25'}[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}NR^{25'}CO$, $NR^{25'}CO[C(R^{24'})_2]_{a''}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{c''}NR^{25'}SO_2$, $CR^{24'}=CR^{24'}CO$, $C\equiv CCO$, $(C(R^{24'})_2)_{c''}SO_2$, $SO_2[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}SO_2$, $O[C(R^{24'})_2]_{a''}SO_2$, $SO_2NR^{25'}[C(R^{24'})_2]_{1-2}$, $[C(R^{24'})_2]_{b''}COO[C(R^{24'})_2]_2$, $[C(R^{24'})_2]_{b''}CONR^{25'}[C(R^{24'})_2]_{1-2}$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$, then D' may further be O, $NR^{25'}$, $CONR^{25'}$, $SO_2NR^{25'}$, $OCONR^{25'}$, $NR^{25'}COO$, $NR^{25'}CONR^{25'}$, $[C(R^{24'})_2]_{a''}NR^{25'}[C(R^{24'})_2]_{b''}$, $[C(R^{24'})_2]_{a''}O[C(R^{24'})_2]_{b''}$, $CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}O$, $NR^{25'}[C(R^{24'})_2]_{a''}NR^{25'}$, $O[C(R^{24'})_2)]_{a''}NR^{25'}$, $O[C(R^{24'})_2]_{a''}O$, $CO[C(R^{24'})_2]_{a''}O$, $SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $SO_2[C(R^{24'})_2]_{a''}O$, $[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $[C(R^{24'})_2]_{a''}CONR^{25'}$, $O[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $O[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $(C(R^{24'})_2)_{a''}S(C(R^{24'})_2)_{b''}$, COO, $CR^{24'}OH$, $C(R^{24'})_{a''}CR^{24'}OH$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$ or $C=CR^{26'}$, D' may further be $CR^{24'}=CR^{24'}$ or $C\equiv C$; and a'' is 1-6, b'' is 0-1, c'' is 0-2;

$R^{24'}$ is hydrogen or C_{1-6}alkyl;

$R^{25'}$ is hydrogen or C_{1-6}alkyl;

E' and G' together are $NC(R^{26'})_2$, $NC(R^{26'})_2C(R^{26'})_2$, $CR^{27'}C(R^{26'})_2$ or $C=CR^{26'}$;

$R^{26'}$ is hydrogen or C_{1-6}alkyl;

$R^{27'}$ is hydrogen, $OR^{28'}$, $NHR^{28'}$, CN, NO_2, $R^{28'}$, $SR^{29'}$, $COR^{28'}$, $CHOHR^{28'}$, $CO_2R^{28'}$, $NHCOR^{28'}$, $NHCO_2R^{29'}$, $NHSO_2R^{29'}$, or $OCONHR^{28'}$;

$R^{28'}$ is hydrogen, C_{1-5}alkyl, aryl or aralkyl;

$R^{29'}$ is C_{1-5}alkyl, aryl or aralkyl;

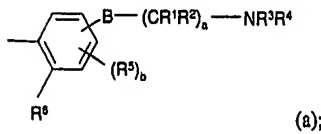
R' is one or more of hydrogen or C_{1-6}alkyl, or R' is oxo;

J' is CO or SO_2;

L' is $NR^{30'}$, O or $C(R^{30'})_2$;

$R^{30'}$ is hydrogen or C_{1-6}alkyl;

E represents a group (a):



(a);

in which

B is oxygen, C≡C, $S(O)_c$, $CR^7=CR^8$, or CR^7R^8, or B is NR^9;

R^1 and R^2 are independently hydrogen or C_{1-6}alkyl; alternatively $B(CR^1R^2)_a$ is $OCR^1R^2CR^1(OH)CR^1R^2$ or $OCR^1R^2CR^1(OCOCH_3)CR^1R^2$;

R^3 and R^4 are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{10}R^{11}$, $NR^{10}R^{11}$, hydroxy, $OCOR^{12}$, $NHCOCF_3$, $NHSO_2R^{13}$, $NHCO_2R^{14}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

R^5 is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{15}R^{16}$, CO_2R^{17}, trifluoromethyl, $NHCO_2R^{18}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_dR^{19}$, $SO_2NR^{20}R^{21}$ or halogen;

R^6 is hydrogen, C_{1-6}alkyl, aryl, trifluoromethyl, hydroxy, C_{1-6}alkoxy or halogen, or R^6 taken together with $R^{30'}$ forms a group D where D is $(CR^{22}R^{23})_e$ or D is $(CR^{22}R^{23})_f$-G where G is oxygen, sulfur or $CR^{22}=CR^{23}$, $CR^{22}=N$, $=CR^{22}O$, $=CR^{22}S$, or $=CR^{22}-NR^{23}$;

R^7, R^8, R^{10}, R^{11}, R^{12}, R^{15}, R^{16}, R^{17}, R^{20}, R^{21}, R^{22}, and R^{23} are independently hydrogen or C_{1-6}alkyl;

R^9 is hydrogen, C_{1-6}alkyl, or phenylC_{1-6}alkyl;

R^{13}, R^{14}, R^{18}, and R^{19} are independently C_{1-6}alkyl;

a is 1, 2, 3, or 4;

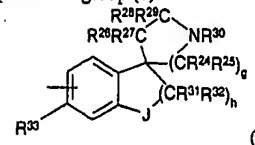
b is 1 or 2;

c and d are independently 0, 1 or 2;

e is 2, 3 or 4;

f is 0, 1, 2 or 3;

alternatively, E represents a group (b):



(b);

R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31}, and R^{32} are independently hydrogen or

C_{1-6}alkyl;

R^{30} is hydrogen, C_{1-6}alkyl, or C_{3-7}cycloalkyl;

R^{33} is hydrogen, C_{1-6}alkyl, trifluoromethyl, hydroxy or halogen, or R^{33} and $R^{30'}$ together form a group -K- where K is $(CR^{34}R^{35})_i$ or K is $(CR^{34}R^{35})_j$ -M and M is oxygen, sulfur, $CR^{34}=CR^{35}$, $CR^{34}=N$, or N=N;

J is oxygen, $CR^{36}R^{37}$, or NR^{38}, or J is a group $S(O)_k$;

R^{34}, R^{35}, R^{36}, R^{37}, and R^{38} are independently hydrogen or C_{1-6}alkyl;

g is 1, 2 or 3;

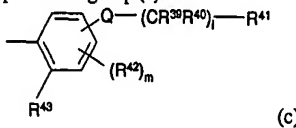
h is 1, 2 or 3;

i is 2, 3, or 4;

j is 0, 1, 2, or 3;

k is 0, 1 or 2;

alternatively, E represents a group (c):



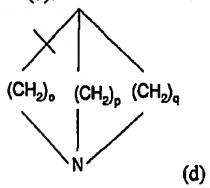
(c);

in which:

Q is oxygen, $S(O)_n$, $CR^{44}=CR^{45}$, $CR^{44}R^{45}$, or Q is NR^{46};

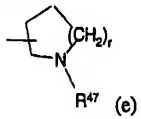
R^{39} and R^{40} are independently hydrogen or C_{1-6}alkyl;

R^{41} is a group of formula (d):



(d)

or R^{41} is a group of formula (e):



(e)

R^{42} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{48}R^{49}$, CO_2R^{50}, trifluoromethyl, $NHCO_2R^{51}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_sR^{52}$, $SO_2NR^{53}R^{54}$, or halogen;

R^{43} is hydrogen or R^{43} together with $R^{30'}$ forms a group R where R is $CR^{55}=CR^{56}$, $CR^{55}=CR^{56}CR^{55}R^{56}$, or $(CR^{55}R^{56})t$;

R^{44}, R^{45}, R^{46}, R^{48}, R^{49}, R^{50}, R^{53}, R^{54}, R^{55}, and R^{56} are independently hydrogen or C_{1-6}alkyl;

R^{47} is hydrogen, C_{1-6}alkyl, or C_{3-7} cycloalkyl;

R^{51} and R^{52} are independently C_{1-6}alkyl;

l is 0, 1, 2, or 3;

m is 1 or 2;

n is 0, 1, or 2

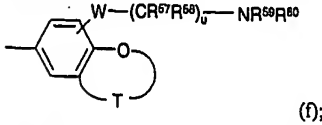
o, p, and q are independently integers having the value 1, 2, or 3;

r is 0,1, 2, or 3;

s is 0, 1, or 2;

t is 2 or 3;

alternatively, E represents a group (f):



(f);

R^{57} and R^{58} are independently hydrogen or C_{1-6}alkyl;

R^{59} and R^{60} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{61}R^{62}$, $NR^{61}R^{62}$, hydroxy, $OCOR^{63}$, $NHCOCF_3$, $NHSO_2R^{64}$, $NHCO_2R^{65}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

T is $-(CR^{66}R^{67})_v-$ or $-O(CR^{66}R^{67})_w-$;

W is oxygen, $S(O)_x$, NR^{68}, or W is $CR^{69}=CR^{70}$ or $CR^{69}R^{70}$;

R^{61}, R^{62}, R^{63}, R^{66}, R^{67} R^{68}, R^{69}, and R^{70} are independently hydrogen or C_{1-6}alkyl;

R^{64} and R^{65} are independently C_{1-6}alkyl;

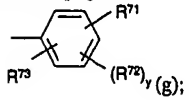
u is 1 to 4;

v is 2 or 3;

w is 1, 2, or 3;

x is 0, 1 or 2;

alternatively, E represents a group (g):



(g);

R^{71} is a 5- to 7-membered saturated or partially saturated heterocyclic ring containing a nitrogen atom and optionally a further 1 or 2 heteroatoms selected from nitrogen, oxygen or sulfur or R^{71} is an optionally substituted 6,6 or 6,5 bicyclic ring containing a nitrogen atom and optionally a further heteroatom selected from oxygen, nitrogen or sulfur, which ring systems may be optionally substituted with one or more of C_{1-6}alkyl and optionally substituted on nitrogen with hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl;

R^{72} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{74}R^{75}$, CO_2R^{76}, trifluoromethyl, $NHCO_2R^{77}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_zR^{78}$, $SO_2NR^{79}R^{80}$, or halogen;

R^{73} is hydrogen, C_{1-6}alkyl, hydroxy, C_{1-6}alkoxy or halogen, or R^{73} and $R^{30'}$ taken together from a group -X- where X is $(CR^{81}R^{82})_{aa}$ or X is $(CR^{81}R^{82})_{ab}$-Y and Y is oxygen, sulfur or $CR^{81}=CR^{82}$;

R^{74}, R^{75}, R^{76}, R^{79}, R^{80}, R^{81}, and R^{82} are independently hydrogen or C_{1-6}alkyl;

R^{77} and R^{78} are independently C_{1-6}alkyl;

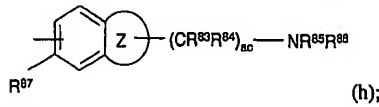
y is 1 or 2;

z is 0, 1, or 2;

aa is 2, 3 or 4;

ab is 0, 1, 2 or 3;

alternatively, E represents a group (h):



(h);

R^{83} and R^{84} are independently hydrogen or C_{1-6}alkyl;

R^{85} and R^{86} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{88}R^{89}$, $NR^{90}R^{91}$, hydroxy, $OCOR^{92}$, $NHCOCF_3$, $NHSO_2R^{93}$, $NHCO_2R^{94}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

R^{87} is hydrogen or C_{1-6}alkyl, C_{1-6}alkoxy, or halogen, or R^{87} together with $R^{30'}$ forms a group -AA- where AA is $(CR^{95}R^{96})_{ad}$ or AA is $(CR^{95}=CR^{96})_{ae}$-AB and AB is oxygen, sulfur, $CR^{95}=CR^{96}$, $CR^{95}=N$, $CR^{95}NR^{96}$ or N=N;

Z is an optionally substituted 5 to 7-membered heterocyclic ring containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulfur;

R^{88}, R^{89}, R^{90}, R^{91}, R^{92}, R^{95}, and R^{96} are independently hydrogen or C_{1-6}alkyl;

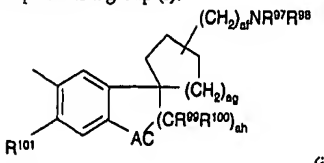
R^{93} and R^{94} are independently C_{1-6}alkyl;

ac is 0 to 4;

ad is 1, 2 or 3;

ae is 0, 1 or 2;

alternatively, E represents a group (i):



(i);

R^{97} and R^{98} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{102}R^{103}$, $NR^{104}R^{105}$, hydroxy, $OCOR^{106}$, $NHCOCF_3$, $NHSO_2R^{107}$, $NHCO_2R^{108}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

R^{99} and R^{100} are independently hydrogen or C_{1-6}alkyl;

R^{101} is hydrogen or C_{1-6}alkyl or R^{101} and $R^{30'}$ together form a group -AD- where AD is $(CR^{109}R^{110})_{ai}$ or AD is $(CR^{109}R^{110})_{aj}$-AE and AE is oxygen, sulfur or $CR^{109}=CR^{110}$;

AC is oxygen, $CR^{111}R^{112}$ or NR^{113} or AC is a group $S(O)_{ak}$;

R^{102}, R^{103}, R^{104}, R^{105}, R^{106}, R^{109}, R^{110}, R^{111}, R^{112}, and R^{113} are independently hydrogen or C_{1-6}alkyl;

R^{107} and R^{108} are independently C_{1-6}alkyl;

af is 0, 1, 2, 3, or 4;

ag is 1, 2, or 3;

ah is 1, 2, 3 or 4;

ai is 2, 3 or 4;

aj is 0, 1, 2, or 3; and

ak is 0, 1 or 2.

For compounds of formula (I) various embodiments are as follows. It will be understood that the basic nitrogen in moiety E may be optionally quaternized with C_{1-6}alkyl or is optionally present as the N-oxide.

Suitably, A' is aryl or heteroaryl, each of which is optionally substituted with

one or more of $R^{1'}$. Alternatively, A' is suitably aryl or heteroaryl fused to a saturated or partly unsaturated 5-7-membered ring to form a higher order ring moiety, which ring moiety optionally contains 1 or 2 heteroatoms selected from oxygen, nitrogen or sulfur, wherein nitrogen may be optionally substituted with hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl; wherein the higher order ring moiety is optionally substituted with one or more of $R^{1'}$. Preferably A' is phenyl, 5,6,7,8-tetrahydro-1-naphthalenyl, 1H-indol-4-yl, or 2-benzothiazolyl.

Suitably, $R^{1'}$ is hydrogen, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, CH_2CF_3, aryl, aralkyl, $(CH_2)_a{}'NR^{2'}R^{3'}$, $(CH_2)_a{}'NR^{2'}COR^{4'}$, $(CH_2)_aNR^{2'}CO_2R^{5'}$, $(CH_2)_a{}'NR^{2'}SO_2R^{6'}$, $(CH_2)_a{}'CONR^{7'}R^{8'}$, hydroxy$C_{1-6}$alkyl, C_{1-4}alkoxyalkyl (optionally substituted by a C_{1-4}alkoxy or hydroxy group), $(CH_2)_a{}'CO_2C_{1-6}$alkyl, $(CH_2)_b{}'OC(O)R^{9'}$, $CR^{10'}=NOR^{11'}$, $CNR^{10'}=NOR^{11'}$, $COR^{12'}$, $CONR^{7'}R^{8'}$, $CONR^{7'}(CH_2)_c{}'OC_{1-4}$alkyl, $CONR^{7'}(CH_2)_a{}'CO_2R^{13'}$, $CONHNR^{14'}R^{15'}$, $CONR^{7'}SO_2R^{16'}$, $CO_2R^{17'}$, cyano, trifluoromethyl, $NR^{2'}R^{3'}$, $NR^{2'}COR^{4'}$, $NR^{18'}CO(CH_2)_a{}'NR^{18'}R^{19'}$, $NR^{18'}CONR^{18'}R^{19'}$, $NR^{2'}CO_2R^{5'}$, $NR^{2'}SO_2R^{6'}$, $N{=}CNR^{18'}NR^{18'}R^{19'}$, nitro, hydroxy, C_{1-6}alkoxy, OCF_3, hydroxyC_{1-6}alkoxy, C_{1-6}alkoxyC_{1-6}alkoxy, $OC(O)NR^{20'}R^{21'}$, $SR^{22'}$, $SOR^{23'}$, $SO_2R^{23'}$, $SO_2NR^{20'}R^{21'}$ or halogen, or suitably $R^{1'}$ is a 5- to 7-membered heterocyclic ring containing 1 to 4 heteroatoms selected from oxygen, nitrogen, or sulfur, suitable heterocyclic rings include aromatic groups such as thienyl, furyl, pyrrolyl, triazolyl, diazolyl, imidazolyl, oxazolyl, thiazolyl, oxadiazolyl, isothiazolyl, isoxazolyl, thiadiazolyl, pyridyl, pyrimidyl, pyrazinyl, and dioxanyl. Saturated and partially saturated rings are also within the scope of the invention, in particular rings including an oxo or thioxo moiety such as lactams and thiolactams. Suitably, the heterocyclic ring can be linked to the remainder of the molecule via a carbon atom, or, when present, a nitrogen atom. Suitably these rings may be optionally substituted with one or more of hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, hydroxyC_{1-6}alkyl, $(C_{1-6}$alkyl$)C_{1-6}$alkyl, $CONR^{7'}R^{8'}$, $CO_2R^{17'}$, cyano, aryl, trifluoromethyl, nitro, hydroxy, C_{1-6}alkoxy, acyloxy, or halogen. Preferably, $R^{1'}$ is one or more of C_{1-6}alkyl, $(CH_2)_aNR^{2'}COR^{4'}$, CF_3, CO_2C_{1-6}alkyl, C_{1-6}alkoxy, halogen, or cyano.

Suitably, a' is 1, 2, 3 or 4; b' is 0, 1, 2 or 3; and c' is 1, 2 or 3.

Suitably, $R^{2'}$ and $R^{3'}$ are independently hydrogen or C_{1-6}alkyl, or suitably, $R^{2'}$ and $R^{3'}$ together with the nitrogen to which they are attached, form a 5- to 6-membered heterocyclic ring. Suitably, the ring may be optionally substituted by an oxo group, or, when $R^{2'}$ and $R^{3'}$ form a 6-membered ring, the ring may optionally contain one oxygen or one sulfur atom. When the ring is a 6-membered ring substituted by an oxygen or

sulfur atom, the oxygen or sulfur atom are preferably in the 4-position.

Suitably, $R^{4'}$ is hydrogen, C_{1-6}alkyl or C_{1-4}alkoxyalkyl, or, when $R^{1'}$ is $NR^{2'}COR^{4'}$, $R^{4'}$ is $(CH_2)_{1-3}$ and forms a ring with A'.

Suitably $R^{5'}$ is C_{1-6}alkyl.

Suitably, $R^{6'}$ is C_{1-6}alkyl or phenyl.

Suitably, $R^{7'}$ and R^{8} are independently hydrogen or C_{1-6}alkyl, or suitably, $R^{7'}$ and $R^{8'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring. Suitably, when the ring is 6-membered, the ring may optionally contain one oxygen or one sulfur atom.

Suitably, $R^{9'}$ is C_{1-4}alkyl, wherein the C_{1-6}alkyl is optionally substituted by a C_{1-6}alkoxy.

Suitably, $R^{10'}$ and $R^{11'}$ are independently hydrogen or C_{1-6}alkyl.

Suitably, $R^{12'}$ is hydrogen or C_{1-6}alkyl.

Suitably, $R^{13'}$ is hydrogen or C_{1-6}alkyl.

Suitably, $R^{14'}$ and $R^{15'}$ are independently hydrogen or C_{1-6}alkyl.

Suitably, $R^{16'}$ is hydrogen or C_{1-6}alkyl.

Suitably, $R^{17'}$ is hydrogen or C_{1-6}alkyl, wherein the C_{1-6}alkyl is optionally substituted with one or more substituents selected from C_{1-6}alkyl, C_{1-6}alkoxy, hydroxy, or $NR^{2'}R^{3'}$. Preferably, when there is more than one substituent, there are two substituents.

Suitably, $R^{18'}$ and $R^{19'}$ are independently hydrogen or C_{1-6}alkyl.

Suitably, $R^{20'}$ and $R^{21'}$ are independently hydrogen or C_{1-6}alkyl, or suitably, $R^{20'}$ and $R^{21'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring which, when there are 6 ring members, may optionally contain in the ring one oxygen or one sulfur atom.

Suitably, $R^{22'}$ is hydrogen or C_{1-6}alkyl.

Suitably, $R^{23'}$ is C_{1-6}alkyl.

Suitably, D' is either a bond or represents $[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{a''}CO$, SO_2, CO, $CO[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}$, $S[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}OCO$, $NR^{25'}[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}NR^{25'}CO$, $NR^{25'}CO[C(R^{24'})_2]_{a''}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{c''}NR^{25'}SO_2$, $CR^{24'}{=}CR^{24'}CO$, $C{\equiv}CCO$, $(C(R^{24'})_2)_{c''}SO_2$, $SO_2[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}SO_2$, $O[C(R^{24'})_2]_{a''}SO_2$, $SO_2NR^{25'}[C(R^{24'})_2]_{1-2}$, $[C(R^{24'})_2]_{b''}COO[C(R^{24'})_2]_2$, $[C(R^{24'})_2]_{b''}CONR^{25'}[C(R^{24'})_2]_{1-2}$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$, then D' may further be O, $NR^{25'}$, $CONR^{25'}$, $SO_2NR^{25'}$, $OCONR^{25'}$, $NR^{25'}COO$, $NR^{25}CONR^{25'}$, $[C(R^{24'})_2]_{a''}NR^{25'}[C(R^{24'})_2]_{b''}$,

$[C(R^{24'})_2]_{a''}O[C(R^{24'})_2]_{b''}$, $CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}O$, $NR^{25'}[C(R^{24'})_2]_{a''}NR^{25'}$, $O[C(R^{24'})_2)]_{a''}NR^{25'}$, $O[C(R^{24'})_2]_{a''}O$, $CO[C(R^{24'})_2]_{a''}O$, $SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $SO_2[C(R^{24'})_2]_{a''}O$, $[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $[C(R^{24'})_2]_{a''}CONR^{25'}$, $O[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $O[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $(C(R^{24'})_2)_{a''}S(C(R^{24'})_2)_{b''}$, COO, $CR^{24'}OH$, $C(R^{24'})_{a''}CR^{24'}OH$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$ or $C{=}CR^{26'}$, D' may further be $CR^{24}{=}CR^{24}$ or $C{\equiv}C$; and a'' is 1-6, b'' is 0-1, c'' is 0-2. Preferably, D' is a bond, CO or SO_2.

Suitably, $R^{24'}$ is hydrogen or C_{1-6}alkyl.

Suitably, $R^{25'}$ is hydrogen or C_{1-6}alkyl.

Suitably, E' and G' together are $NC(R^{26'})_2$, $NC(R^{26'})_2C(R^{26'})_2$, $CR^{27'}C(R^{26'})_2$ or $C{=}CR^{26'}$. Preferably, E' and G' together are $NC(R^{26'})_2$.

Suitably, $R^{26'}$ is hydrogen or C_{1-6}alkyl. Preferably, $R^{26'}$ is hydrogen.

Suitably, $R^{27'}$ is hydrogen, $OR^{28'}$, $NHR^{28'}$, CN, NO_2, $R^{28'}$, $SR^{29'}$, $COR^{29'}$, $CHOHR^{29'}$, $CO_2R^{29'}$, $NHCOR^{29'}$, $NHCO_2R^{29'}$, $NHSO_2R^{29'}$, or $OCONHR^{29'}$.

Suitably, $R^{28'}$ is hydrogen, C_{1-5}alkyl, aryl or aralkyl.

Suitably, $R^{29'}$ is C_{1-5}alkyl, aryl or aralkyl.

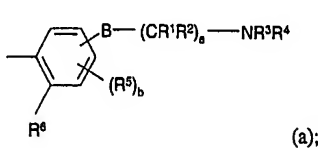
Suitably, R' is one or more of hydrogen or C_{1-6}alkyl, or R' is oxo. Preferably, R' is hydrogen.

Suitably, J' is CO or SO_2. Preferably, J' is CO.

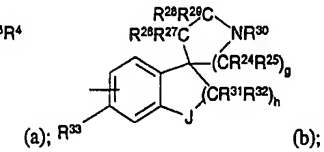
Suitably, L' is $NR^{30'}$, O, or $C(R^{30'})_2$. Preferably, L' is $NR^{30'}$.

Suitably, $R^{30'}$ is hydrogen or C_{1-6}alkyl. Preferably, $R^{30'}$ is hydrogen.

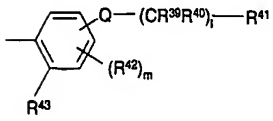
Suitably, substituent E is selected from the following groups:



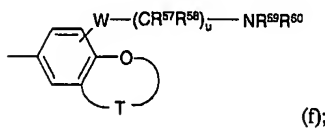
(a);



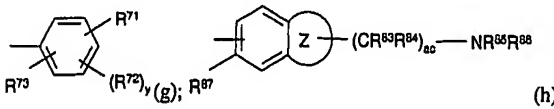
(b);



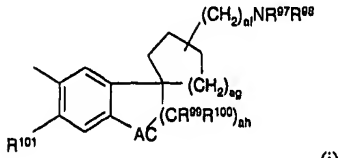
(c);



(f);

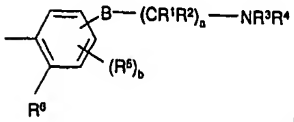


(h); and



(i).

E suitably represents a group (a):



(a).

B is suitably oxygen, C≡C, $S(O)_c$, $CR^7{=}CR^8$, or CR^7R^8, or B is NR^9. B is preferably CR^7R^8, or oxygen.

R^1 and R^2 are suitably independently hydrogen or C_{1-6}alkyl. Preferably, R^1 and R^2 are each hydrogen. Alternatively, $B(CR^1R^2)_a$ is $OCR^1R^2CR^1(OH)CR^1R^2$ or $OCR^1R^2CR^1(OCOCH_3)CR^1R^2$. Preferably, when $B(CR^1R^2)_a$ is $OCR^1R^2CR^1(OH)CR^1R^2$ or $OCR^1R^2CR^1(OCOCH_3)CR^1R^2$, R^1 and R^2 are hydrogen.

R^3 and R^4 are suitably independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{10}R^{11}$, $NR^{10}R^{11}$, hydroxy, $OCOR^{12}$, $NHCOCF_3$, $NHSO_2R^{13}$, $NHCO_2R^{14}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH. Preferably R^3 and R^4 are independently C_{1-6}alkyl, C_{3-7}cycloalkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur.

Preferably, $B\text{-}(CR^1R^2)_a\text{-}NR^3R^4$ is ortho to R^5, meta to L' and para to R^6, and R^5 is para to L'.

R^5 is suitably hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{15}R^{16}$, CO_2R^{17}, trifluoromethyl, $NHCO_2R^{18}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_dR^{19}$, $SO_2NR^{20}R^{21}$, or halogen. R^5 is preferably C_{1-6}alkoxy, SC_{1-6}alkyl or halogen.

R^6 is suitably hydrogen, C_{1-6}alkyl, aryl, trifluoromethyl, hydroxy, C_{1-6}alkoxy, or halogen, or R^6 taken together with $R^{30'}$ forms a group D where D is $(CR^{22}R^{23})_e$ or D is $(CR^{22}R^{23})_f$-G where G is oxygen, sulfur, or $CR^{22}{=}CR^{23}$, $CR^{22}{=}N$, $=CR^{22}O$, $=CR^{22}S$, or $=CR^{22}\text{-}NR^{23}$. Preferably, R^6 is hydrogen.

R^7, R^8, R^{10}, R^{11}, R^{12}, R^{15}, R^{16}, R^{17}, R^{20}, R^{21}, R^{22}, and R^{23} are suitably independently hydrogen or C_{1-6}alkyl.

R^9 is suitably hydrogen, C_{1-6}alkyl, or phenylC_{1-6}alkyl.

R^{13}, R^{14}, R^{18}, and R^{19} are suitably independently C_{1-6}alkyl.

a is suitably 1, 2, 3, or 4. Preferably, a is 2 or 3.

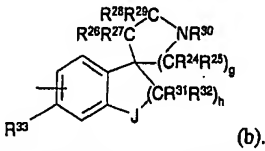
b is suitably 1 or 2. Preferably, b is 1.

c and d are suitably independently 0, 1, or 2.

e is suitably 2, 3, or 4.

f is suitably 0, 1, 2, or 3.

Alternatively, E suitably represents a group (b):



(b).

R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31}, and R^{32} are suitably independently hydrogen or C_{1-6}alkyl. R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31}, and R^{32} are preferably hydrogen.

R^{30} is suitably hydrogen, C_{1-6}alkyl, or C_{3-7}cycloalkyl. Preferably, R^{30} is C_{1-6}alkyl or C_{3-7}cycloalkyl.

R^{33} is suitably hydrogen, C_{1-6}alkyl, trifluoromethyl, hydroxy or halogen, or R^{33} and $R^{30'}$ together form a group -K- where K is $(CR^{34}R^{35})_i$ or K is $(CR^{34}R^{35})_j$-M and M is oxygen, sulfur, $CR^{34}{=}CR^{35}$, $CR^{34}{=}N$, or N=N. Preferably, R^{33} is hydrogen.

J is suitably oxygen, $CR^{36}R^{37}$, or NR^{38}, or J is a group $S(O)_k$. Preferably, J is oxygen. Preferably, J is para to L'.

R^{34}, R^{35}, R^{36}, R^{37}, R^{38} are suitably independently hydrogen or C_{1-6}alkyl.

g is suitably 1, 2, or 3. Preferably, g is 2 or 3.

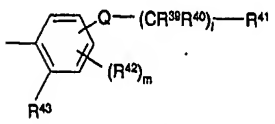
h is suitably 1, 2, or 3. Preferably, h is 1.

i is suitably 2, 3, or 4.

j is suitably 0, 1, 2, or 3.

k is suitably 0, 1 or 2.

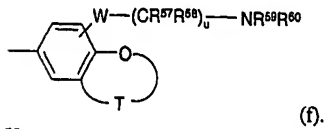
Alternatively, E suitably represents a group (c):



(c).

Suitably, Q is oxygen, $S(O)_n$, $CR^{44}=CR^{45}$, C=C , or $CR^{44}R^{45}$, wherein n is 0, 1 or 2, and R^{44} and R^{45} are independently hydrogen or C_{1-6}alkyl, or suitably, Q is NR^{46} wherein R^{46} is hydrogen or alkyl; suitably, R^{39} and R^{40} are independently hydrogen or C_{1-6}alkyl; suitably, R^{42} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{48}R^{49}$, CO_2R^{50}, trifluoromethyl, $NHCO_2R^{51}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_8R^{52}$, $SO_2NR^{53}R^{54}$, or halogen, wherein R^{48}, R^{49}, R^{50}, R^{53}, and R^{54} are hydrogen or C_{1-6}alkyl, and R^{51} and R^{52} are C_{1-6}alkyl; suitably, R^{43} is hydrogen or R^{43} together with $R^{30'}$ forms a group R where R is $CR^{55}=CR^{56}$, $CR^{55}=CR^{56}CR^{55}R^{56}$, or $(CR^{55}R^{56})_t$ wherein R^{55} and R^{56} are independently hydrogen or C_{1-6}alkyl and t is 2 or 3; suitably, R^{41} is selected from a group of formula (d) or (e); suitably R^{47} is hydrogen, C_{1-6}alkyl, or C_{3-7} cycloalkyl; suitably, l is 0, 1, 2 or 3, m is 1 or 2, n and s are independently 0, 1 or 2, o, p and q are independently 1, 2 or 3, and r is 0, 1, 2 or 3.

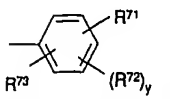
Alternatively, E suitably represents a group (f):



(f).

Suitably, R^{57} and R^{58} are independently hydrogen or C_{1-6}alkyl; suitably R^{59} and R^{60} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{61}R^{62}$, $NR^{61}R^{62}$, hydroxy, $OCOR^{63}$, $NHCOCF_3$, $NHSO_2R^{64}$, $NHCO_2R^{65}$ or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH, and wherein R^{61}, R^{62}, and R^{63} are independently hydrogen or C_{1-6}alkyl, and R^{64} and R^{65} are independently C_{1-6}alkyl; suitably, T is $-(CR^{66}R^{67})_{v}-$ or $-O(CR^{66}R^{67})_{w}-$, wherein R^{66} and R^{67} are independently hydrogen or C_{1-6}alkyl, wherein v is 2 or 3, and w is 1, 2 or 3; suitably, W is oxygen, $S(O)_x$, wherein x is 0, 1 or 2, or W is NR^{68}, wherein R^{68} is hydrogen or C_{1-6}alkyl, or W is $CR^{69}=CR^{70}$, C=C, or $CR^{69}R^{70}$, wherein R^{69} and R^{70} are independently hydrogen or C_{1-6}alkyl; and suitably, u is an integer from 1-4.

Alternatively, E suitably represents a group (g):



(g).

Suitably, R^{71} is an optionally substituted 5- to 7-membered saturated or partially saturated heterocyclic ring containing a nitrogen atom and optionally a further one or two heteroatoms selected from nitrogen, oxygen or sulfur, or R^{71} is an optionally substituted 6,6 or 6,5-bicyclic ring system containing a nitrogen atom and optionally a further heteroatom selected from oxygen, nitrogen or sulfur, which ring systems may be optionally substituted with one or more of C_{1-6}alkyl, and substituted on nitrogen with hydrogen, C_{1-6}alkyl, or C_{3-7}cycloalkyl. Examples of such ring systems include, but are not limited to, pyrrolidine, piperidine, piperazine, morpholine, imidazolidine, pyrazolidine, 1,2,3,6-tetrahydropyridine, hexahydroazepine, tropane, isoquinuclidine and granatane rings. Preferably, R^{71} is an optionally substituted 5- or 6-membered saturated or partially saturated heterocyclic ring containing a nitrogen atom and substituted on nitrogen with C_{1-6}alkyl or C_{3-7}cycloalkyl.

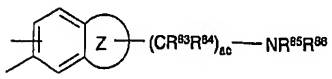
R^{71} is preferably located meta to L', ortho to R^{72} and para to R^{73}, and R^{72} is located para to L'.

Suitably, R^{72} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{74}R^{75}$, CO_2R^{76}, trifluoromethyl, $NHCO_2R^{77}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_zR^{78}$, $SO_2NR^{79}R^{80}$, or halogen wherein R^{74}, R^{75}, R^{76}, R^{79} and R^{80} are independently hydrogen or C_{1-6}alkyl, R^{77} and R^{78} are C_{1-6}alkyl, and z is 0, 1, or 2. R^{72} is preferably C_{1-6}alkoxy, SC_{1-6}alkyl or halogen.

R^{73} is hydrogen, C_{1-6}alkyl, hydroxy, C_{1-6}alkoxy or halogen, or R^{73} and $R^{30'}$ taken together from a group -X- where X is $(CR^{81}R^{82})_{aa}$, wherein aa is 2, 3 or 4, and R^{81} and R^{82} are independently hydrogen or C_{1-6}alkyl, or X is $(CR^{81}R^{82})_{ab}$-Y, wherein ab is 0, 1, 2 or 3, and Y is oxygen, sulfur or $CR^{81}=CR^{82}$ wherein R^{81} and R^{82} are independently hydrogen or C_{1-6}alkyl. Preferably, R^{73} is hydrogen.

Suitably, y is an integer from 1-2. Preferably, y is 1.

Alternatively, E suitably represents a group (h):

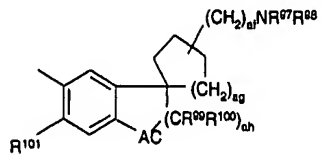


(h).

Suitably, R^{87} is hydrogen, C_{1-6}alkyl, C_{1-6}alkoxy or halogen, or R^{87} together with $R^{30'}$ form a group -AA-, wherein AA is $(CR^{95}R^{88})$ad, wherein ad is 1, 2 or 3, and

R^{95} and R^{88} are independently hydrogen or C_{1-6}alkyl, or AA is $(CR^{95}CR^{96})_{ae}$-AB, wherein ae is 0, 1 or 2, and AB is oxygen, sulfur, $CR^{95}=CR^{96}$, $CR^{95}=N$, $CR^{95}NR^{96}$ or N=N wherein R^{95} and R^{96} are independently hydrogen or C_{1-6}alkyl; suitably, R^{83} and R^{84} are independently hydrogen or C_{1-6}alkyl; suitably, R^{85} and R^{86} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{88}R^{89}$, $NR^{90}R^{91}$, hydroxy, $OCOR^{92}$, $NHCOCF_3$, $NHSO_2R^{93}$, $NHCO_2R^{94}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of the $NHCOC_{0-6}$alkyl is optionally substituted by OH, and wherein R^{88}, R^{89}, R^{90}, R^{91} and R^{92} are independently hydrogen or C_{1-6}alkyl, and R^{93} and R^{94} are independently C_{1-6}alkyl; suitably Z is an optionally substituted 5 to 7-membered heterocyclic ring containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulfur; suitably ac is 0-4.

Alternatively, E suitably represents a group (i):



(i).

Suitably, R^{101} is hydrogen or C_{1-6}alkyl or R^{101} and $R^{30'}$ together form a group -AD- wherein AD is $(CR^{109}R^{110})_{ai}$ wherein ai is 2, 3 or 4 or AD is $(CR^{109}R^{110})_{aj}$-AE wherein aj is 0, 1, 2 or 3 and AE is oxygen, sulfur or $CR^{109}=CR^{110}$, and R^{109} and R^{110} are independently hydrogen or C_{1-6}alkyl; suitably, R^{97} and R^{98} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{102}R^{103}$, $NR^{104}R^{105}$, hydroxy, $OCOR^{106}$, $NHCOCF_3$, $NHSO_2R^{107}$, $NHCO_2R^{108}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH, and wherein R^{102}, R^{103}, R^{104}, R^{105} and R^{106} are independently hydrogen or C_{1-6}alkyl, and R^{107} and R^{108} are independently C_{1-6}alkyl; suitably, R^{99} and R^{100} are independently hydrogen or C_{1-6}alkyl; suitably, AC is oxygen, $CR^{111}R^{112}$ or NR^{113} wherein R^{111}, R^{112} and R^{113} are independently hydrogen or C_{1-6}alkyl or AC is a group S(O)ak wherein ak is 0, 1 or 2; suitably, ag is an integer from 1-3, ah is an integer from 1-4, and af is 0-4.

Suitably, A' is phenyl, 5,6,7,8-tetrahydro-1-naphthalenyl, 1H-indol-4-yl, or 2-benzothiazolyl, $R^{1'}$ is one or more of C_{1-6}alkyl, $(CH_2)_aNR^2COR^4$, CF_3, CO_2C_{1-6}alkyl, C_{1-6}alkoxy, halogen, or cyano, D' is a bond, E' and G' together are $NC(R^{26})_2$, R' is hydrogen, J' is CO, L' is NR^{30}, and E is group (a), (b), (c), (f), (g), (h), or (i).

More preferably, A' is phenyl, 5,6,7,8-tetrahydro-1-napthalenyl, 1H-indol-4-yl, or 6-chloro-2-benzothiazolyl; and when A' is phenyl, $R^{1'}$ is one or more of C_{1-6}alkyl, CF_3, $CO_2CH_2CH_3$, C_{1-6}alkoxy, halogen, or cyano substituted at the 2,3-, 2,4-, 2,5-, 2-, 3-, 4-, 3,4-, and 3,5- positions, D' is a bond, E' and G' together are NCH_2, R' is hydrogen, J' is CO, L' is NH, and E is a group (a), (b), or (g).

Preferably, E is selected from group (a), (b) and (g).

More preferably, when E is group (a), L' is attached to group (a) meta to B-$(CR^1R^2)_a$-NR^3R^4 and para to $(R^5)_b$, wherein B is oxygen or CR^7R^8, R^1 and R^2 are hydrogen, R^5 is methoxy, methylthio or iodo, R^3 and R^4 are independently C_{3-6}alkyl, or R^3 and R^4 taken together with the nitrogen to which they are attached form a 5- or 6-membered heterocyclic ring optionally substituted with one or more of C_{1-6}alkyl and acetamido or hydroxyl, R^6 is hydrogen, a is 2 or 3 when B is oxygen and a is 2 when B is CR^7R^8, and b is 1.

Most preferably, when E is group (a), L' is attached to group (a) meta to B-$(CR^1R^2)_a$-NR^3R^4 and para to $(R^5)_b$, wherein B is oxygen or CH_2, R^1 and R^2 are hydrogen, R^5 is methoxy, R^3 and R^4 are independently isopropyl or tert-butyl, or R^3 and R^4 taken together with the nitrogen to which they are attached are 1-(2,2,6,6-tetramethylpiperidinyl), 1-(4-acetamido-2,2,6,6-tetramethyl piperidinyl), 1-(4-hydroxy-2,2,6,6-tetramethyl piperidinyl) or 1-(4-hydroxy-2,2,4,6,6-pentamethyl piperidinyl), R^6 is hydrogen, a is 2 when B is oxygen, and b is 1.

More preferably, when E is group (b), L' is attached to group (b) para to J, J is oxygen, R^{33} is hydrogen, R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31} and R^{32} are hydrogen, R^{30} is C_{3-6}alkyl, g is 2 and h is 1.

Most preferably, when E is group (b), L' is attached to group (b) para to J, J is oxygen, R^{33} is hydrogen, R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31} and R^{32} are hydrogen, R^{30} is isopropyl, g is 2, and h is 1.

More preferably, when E is group (g), L' is attached to group (g) meta to R^{71} and para to R^{72}, R^{71} is an optionally substituted 5- or 6-membered saturated or partially saturated heterocyclic ring containing a nitrogen atom substituted on nitrogen with C_{3-6}alkyl or C_{3-7}cycloalkyl, R^{72} is methoxy, methylthio or iodo, y is 1, and R^{73} is hydrogen.

Most preferably, when E is group (g), L' is attached to group (g) meta to R^{71} and para to R^{72} wherein R^{71} is piperidin-4-yl substituted on nitrogen with isopropyl,

R^{72} is methoxy, y is 1, and R^{73} is hydrogen.

A particularly effective subgenus of compounds of formula (I) is wherein, A' is phenyl, 5,6,7,8-tetrahydro-1-naphthalenyl, or 1H-indol-4-yl; and when A' is phenyl, $R^{1'}$ is methyl, chloro or trifluoromethyl substituted at the 2 and/or 3-positions, or $R^{1'}$ is 2,4-dimethyl, 2-methoxy-5-chloro, 2-methyl, 3-ethoxycarbonyl, or 3,5-dichloro, D' is a bond, E' and G' together are NCH_2, R' is hydrogen, J' is CO, L' is NH, and E is group (g).

The term "acyloxy" is used herein at all occurrences to mean a moiety -O-C(O)-R, wherein R is hydrogen or C_{1-6}alkyl as defined below.

The term " C_{1-4}alkanoyl " is used herein at all occurrences to mean a -C(O)C_{1-4}alkyl group wherein the alkyl portion is as defined below.

The term "alkenyl" is used herein at all occurrences to mean a straight or branched chain radical of 2 to 6 carbon atoms, unless the length is limited thereto, wherein there is at least one double bond between two of the carbon atoms in the chain, including, but not limited to, ethenyl, 1-propenyl, 2-propenyl, 2-methyl-1-propenyl, 1-butenyl, 2-butenyl, and the like.

The term "alkoxy" is used herein at all occurrences to mean a straight or branched chain radical of 1 to 6 carbon atoms, unless the chain length is limited thereto, bonded to an oxygen atom, including, but not limited to, methoxy, ethoxy, n- propoxy, isopropoxy, and the like.

The term "C_{1-6}alkoxyC_{1-6}alkoxy" is used herein at all occurrences to mean an alkoxy group as defined above, substituted with an alkoxy group as defined above.

The term "C_{1-4}alkoxyalkyl" is used herein at all occurrences to mean a C_{1-4}alkoxy group as defined above bonded to an alkyl group as defined below, including, but not limited to, -CH_2-CH_2-O-CH_2-CH_2-CH_3 and the like.

The term "C_{1-6}alkyl" is used herein at all occurrences to mean a straight or branched chain radical of 1 to 6 carbon atoms, unless the chain length is limited thereto, including, but not limited to, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, and the like.

The term "alkynyl" is used herein at all occurrences to mean a straight or branched chain radical of 2 to 8 carbon atoms, unless the chain length is limited thereto, wherein there is at least one triple bond between two of the carbon atoms in the chain, including, but not limited to, acetylene, 1- propylene, 2-propylene, and the like.

The term "aralkyl" is used herein at all occurrences to mean an aryl moiety as defined above, which is connected to an alkyl moiety as defined below including, but not limited to, benzyl or phenethyl, and the like.

The term "aryl" is used herein at all occurrences to mean a 6-14-membered

substituted or unsubstituted aromatic ring(s) or ring systems which may include bi- or tri-cyclic systems, including, but not limited to, phenyl, naphthalenyl, biphenyl, phenanthryl, anthracenyl, and the like.

The term "6,6 or 6,5 bicyclic ring" is used herein at all occurrences to mean a 6,6 or 6,5-bicyclic ring system containing a nitrogen atom and optionally a further heteroatom selected from nitrogen, oxygen, or sulfur, which ring system may be optionally substituted with C_{1-6}alkyl. Examples of such ring systems include, but are not limited to, tropane, isoquinuclidine and granatane rings.

The term "cycloalkenyl" is used herein at all occurrences to mean cyclic radicals, preferably of 5 to 8 carbons, which have at least one double bond between two of the carbon atoms in the ring, including but not limited to, cyclopentenyl, cyclohexenyl, and the like.

The terms "cycloalkyl" and "cyclic alkyl" are used herein at all occurrences to mean cyclic radicals, preferably comprising 3 to 7 carbon atoms which may be mono- or bicyclo- fused ring systems which may additionally include unsaturation, including, but not limited to, cyclopropyl, cyclopentyl, cyclohexyl, 1,2,3,4-tetrahydronaphthalenyl, and the like.

The terms "halo" or "halogen" are used interchangeably herein at all occurrences to mean radicals derived from the elements chlorine, fluorine, iodine and bromine.

The term "heteroaryl" is used herein at all occurrences to mean a 5-14-membered substituted or unsubstituted aromatic ring(s) or ring systems which may include bi- or tri-cyclic systems, which ring or ring systems contain 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulfur, including, but not limited to, indolyl, benzofuranyl, thianaphthenyl, quinolyl, isoquinolyl, pyrrolyl, furanyl, thienyl, pyridyl, and the like.

The term "hydroxyC_{1-6}alkoxy" is used herein at all occurrences to mean an hydroxyl group bonded to an alkoxy group as defined above including, but not limited to, -O-CH_2-CH(OH)CH_3 and the like.

The terms "hydroxyC_{1-6}alkyl" and "hydroxyalkyl" are used herein interchangeably to mean an hydroxyl group bonded to a C_{1-6}alkyl group as defined above, including, but not limited to, methanol, ethanol, n-propanol, isopropanol, n-butanol, sec-butanol, isobutanol, tert-butanol, and the like.

The term "heterocyclic ring" is used herein at all occurrences to mean a saturated or partially saturated 5-10-membered ring system (unless the cyclic ring system is otherwise limited) in which the ring system contains one to 3 heteroatoms selected from oxygen, sulfur, or nitrogen, which ring system may be optionally substituted with C_{1-6}alkyl. Examples of

such rings include, but are not limited to, piperidine, tetrahydropyridine, piperazine, pyrrolidine, morpholine, imidazolidine, pyrazolidine, hexahydroazepine, and the like. When the heterocyclic ring is fused to a phenyl group, as when E is the group (h), the term "heterocyclic ring", together with the phenyl ring to which it is fused, forms a ring which includes, but is not limited to, dihydro-1,4-benzoxazine and 1,2,3,4-tetrahydroquinoline, which may be optionally substituted by C_{1-6}alkyl or oxo.

The term "heteroatom" is used herein at all occurrences to mean an oxygen atom, a sulfur atom or a nitrogen atom. It will be recognized that when the heteroatom is nitrogen, it may form an NR_a or NR_aR_b moiety, wherein R_a and R_b are, independently, hydrogen or C_1 to C_6 alkyl, or together with the nitrogen to which they are bound, form a saturated or unsaturated 5-, 6- or 7-membered ring, including, but not limited to, pyrrolidine, piperidine, piperazine, morpholine, pyridine, and the like. It will be recognized that the saturated or unsaturated 5-, 6- or 7-membered ring may optionally have one or more additional heteroatoms in the ring.

The term "optionally substituted" is used herein at all occurrences to mean an optionally substituted 5- to 7-membered heterocyclic ring wherein the optional substituents are one or more of C_{1-6}alkyl.

The term "oxo" is used herein at all occurrences to mean a double bonded oxygen atom attached to a chemical moiety as a substituent.

The term "CCR5 mediated disease state" is used herein at all occurrences to mean any disease state which is mediated (or modulated) by CCR5.

Suitably, pharmaceutically acceptable salts of formula (I) include, but are not limited to, salts with inorganic acids such as hydrochloride, sulfate, phosphate, diphosphate, hydrobromide, and nitrate, or salts with an organic acid such as malate, maleate, fumarate, tartrate, succinate, citrate, acetate, lactate, methanesulfonate, p-toluenesulfonate, palmitate, salicylate, and stearate.

The compounds of the invention can exist in unsolvated as well as solvated forms, including hydrated forms. In general, the solvated forms, with pharmaceutically acceptable solvents such as water, ethanol, and the like, are equivalent to the unsolvated forms for purposes of this invention.

The compounds of the present invention may contain one or more asymmetric carbon atoms and may exist in racemic and optically active forms. The stereocenters may be of any combination of R and S configuration, for example, (R,R), (R,S), (S,S) or (S,R). All of these compounds are within the scope of the present invention.

Among the preferred compounds of the invention are the following compounds:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenyl-1,2,3,6-tetrahydropyridine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-acetamido-methylphenyl) piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-trifluoromethylphenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,6-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dihydro-2(1H)-quinolinone-6-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-

cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[2,3-dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-chloro-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1-naphthalenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1H-indol-4-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)-3-methylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-hydroxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide;

4-(2,3-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,3-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethoxyphenyl)-1-piperazinecarboxamide;

4-(2-Benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(2-Benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-4-hydroxy-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-hydroxy-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-

methoxypheny]-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxypheny]-4-(4-hydroxyphenyl)-1-piperidinecarboxamide;

4-(4-Hydroxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperazinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

4-(3,4-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,6-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-

piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-carboxyphenyl)piperazine-1-carboxamide.

Among the more preferred compounds of the invention are the following compounds:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[2,3-Dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-

piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[4-cyano-1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide.

Among the most preferred compounds of the invention are the following compounds:

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide.

Formulation of Pharmaceutical Compositions

The pharmaceutically effective compounds of this invention (and the pharmaceutically acceptable salts thereof) are administered in conventional dosage forms prepared by combining a compound of formula (I) ("active ingredient") in an amount sufficient to treat COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, ("CCR5-mediated disease states") with standard pharmaceutical carriers or diluents according to conventional procedures well known in the art. These procedures may involve mixing, granulating and compressing or dissolving the ingredients as appropriate to the desired preparation.

The pharmaceutical carrier employed may be, for example, either a solid or liquid. Exemplary of solid carriers are lactose, terra alba, sucrose, talc, gelatin, agar, pectin, acacia, magnesium stearate, stearic acid and the like. Exemplary of liquid carriers are syrup, peanut oil, olive oil, water and the like. Similarly, the carrier or diluent may include time delay material well known to the art, such as glyceryl

monostearate or glyceryl distearate alone or with a wax.

A wide variety of pharmaceutical forms can be employed. Thus, if a solid carrier is used, the preparation can be tableted, placed in a hard gelatin capsule in powder or pellet form or in the form of a troche or lozenge. The amount of solid carrier will vary widely but preferably will be from about 25 mg to about 1000 mg. When a liquid carrier is used, the preparation will be in the form of a syrup, emulsion, soft gelatin capsule, sterile injectable liquid such as an ampule or nonaqueous liquid suspension.

The active ingredient may also be administered topically to a mammal in need of treatment or prophylaxis of CCR5 mediated disease states. The amount of active ingredient required for therapeutic effect on topical administration will, of course, vary with the compound chosen, the nature and severity of the disease state being treated and the mammal undergoing treatment, and is ultimately at the discretion of the physician. A suitable dose of an active ingredient is 1.5 mg to 500 mg for topical administration, the most preferred dosage being 1 mg to 100 mg, for example 5 to 25 mg administered two or three times daily.

By topical administration is meant non-systemic administration and includes the application of the active ingredient externally to the epidermis, to the buccal cavity and instillation of such a compound into the ear, eye and nose, and where the compound does not significantly enter the blood stream. By systemic administration is meant oral, intravenous, intraperitoneal and intramuscular administration.

While it is possible for an active ingredient to be administered alone as the raw chemical, it is preferable to present it as a pharmaceutical formulation. The active ingredient may comprise, for topical administration, from 0.001% to 10% w/w, e.g. from 1% to 2% by weight of the formulation although it may comprise as much as 10% w/w but preferably not in excess of 5% w/w and more preferably from 0.1% to 1% w/w of the formulation.

The topical formulations of the present invention, both for veterinary and for human medical use, comprise an active ingredient together with one or more acceptable carrier(s) therefor and optionally any other therapeutic ingredient(s). The carrier(s) must be 'acceptable' in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

Formulations suitable for topical administration include liquid or semi-liquid preparations suitable for penetration through the skin to the site of inflammation such as liniments, lotions, creams, ointments or pastes, and drops suitable for administration to the eye, ear or nose.

Drops according to the present invention may comprise sterile aqueous or oily

solutions or suspensions and may be prepared by dissolving the active ingredient in a suitable aqueous or alcoholic solution of a bactericidal and/or fungicidal agent and/or any other suitable preservative, and preferably including a surface active agent. The resulting solution may then be clarified by filtration, transferred to a suitable container which is then sealed and sterilized by autoclaving or maintaining at 98-100°C for half an hour. Alternatively, the solution may be sterilized by filtration and transferred to the container by an aseptic technique. Examples of bactericidal and fungicidal agents suitable for inclusion in the drops are phenylmercuric nitrate or acetate (0.002%), benzalkonium chloride (0.01%) and chlorhexidine acetate (0.01%). Suitable solvents for the preparation of an oily solution include glycerol, diluted alcohol and propylene glycol.

Lotions according to the present invention include those suitable for application to the skin or eye. An eye lotion may comprise a sterile aqueous solution optionally containing a bactericide and may be prepared by methods similar to those for the preparation of drops. Lotions or liniments for application to the skin may also include an agent to hasten drying and to cool the skin, such as an alcohol or acetone, and/or a moisturizer such as glycerol or an oil such as castor oil or arachis oil.

Creams, ointments or pastes according to the present invention are semi-solid formulations of the active ingredient for external application. They may be made by mixing the active ingredient in finely-divided or powdered form, alone or in solution or suspension in an aqueous or non-aqueous fluid, with the aid of suitable machinery, with a greasy or non-greasy basis. The basis may comprise hydrocarbons such as hard, soft or liquid paraffin, glycerol, beeswax, a metallic soap; a mucilage; an oil of natural origin such as almond, corn, arachis, castor or olive oil; wool fat or its derivatives, or a fatty acid such as stearic or oleic acid together with an alcohol such as propylene glycol. The formulation may incorporate any suitable surface active agent such as an anionic, cationic or non-ionic surfactant such as esters or polyoxyethylene derivatives thereof. Suspending agents such as natural gums, cellulose derivatives or inorganic materials such as silicaceous silicas, and other ingredients such as lanolin, may also be included.

The active ingredient may also be administered by inhalation. By "inhalation" is meant intranasal and oral inhalation administration. Appropriate dosage forms for such administration, such as an aerosol formulation or a metered dose inhaler, may be prepared by conventional techniques. The daily dosage amount of the active ingredient administered by inhalation is from about 0.1 mg to about 100 mg per day, preferably about 1 mg to about 10 mg per day.

In one aspect, this invention relates to a method of treating COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases,

atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, all in mammals, preferably humans, which comprises administering to such mammal an effective amount of a CCR5 receptor modulator, in particular, a compound as depicted in formula (I).

By the term "treating" is meant either prophylactic or therapeutic therapy. Such formula (I) compound can be administered to such mammal in a conventional dosage form prepared by combining the formula (I) compound with a conventional pharmaceutically acceptable carrier or diluent according to known techniques. It will be recognized by one of skill in the art that the form and character of the pharmaceutically acceptable carrier or diluent is dictated by the amount of active ingredient with which it is to be combined, the route of administration and other well-known variables. The formula (I) compound is administered to a mammal in need of treatment for COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection, in an amount sufficient to decrease symptoms associated with these disease states. The route of administration may be oral or parenteral.

In another aspect, the invention relates to a method for modulating factors which exacerbate the symptoms of the CCR5-mediated diseases described herein.

The term parenteral as used herein includes intravenous, intramuscular, subcutaneous, intra-rectal, intravaginal or intraperitoneal administration. The subcutaneous and intramuscular forms of parenteral administration are generally preferred. The daily parenteral dosage regimen will preferably be from about 30 mg to about 300 mg per day of active ingredient. The daily oral dosage regimen will preferably be from about 100 mg to about 2000 mg per day of active ingredient.

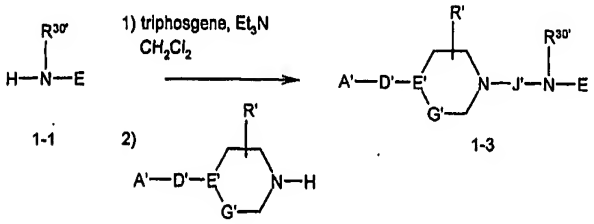
It will be recognized by one of skill in the art that the optimal quantity and spacing of individual dosages of a formula (I) compound will be determined by the nature and extent of the condition being treated, the form, route and site of administration, and the particular mammal being treated, and that such optimums can be determined by conventional techniques. It will also be appreciated by one of skill in the art that the optimal course of treatment, i.e., the number of doses of the formula (I) compound given per day for a defined number of days, can be ascertained by those skilled in the art using conventional course of treatment determination tests.

Methods of Preparation

The compounds of formula (I) can be prepared by art-recognized procedures from known or commercially available starting materials. If the starting materials are unavailable from a commercial source, their synthesis is described herein, or they can be prepared by procedures known in the art.

For example, as shown in Scheme 1, compounds of formula (I) where L' is $NR^{30'}$ are prepared by treating a suitably substituted aniline **1-1** with suitable reagent, for example triphosgene, and a suitable base, for example triethylamine, in a suitable solvent, for example dichloromethane, followed by treatment with a suitably substituted amine **1-2**, e.g., 1-(5,6,7,8-tetrahydro-1-naphthalenyl)piperazine, ethyl 3-(1-piperazinyl)benzoate, 4-(phenyl)piperidine, 1-(phenyl)piperazine, 4-phenyl-2,3,4,6-tetrahydropyrdine, hexahydro-1-phenyl-1H-1,4-diazepine, etc., to afford the title compound **1-3**.

Scheme 1



Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (a) are prepared according to the methods of international application publication number WO 95/15954, published 15 June 1995, international application publication number WO 95/17398, published 29 June 1995, international application publication number WO 95/26328, published 5 October 1995, and international application publication number WO 96/06079, published 29 February 1996.

Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (b) are prepared according to the methods of international application publication number WO 95/11934, published 25 April 1995, and WO 95/19477, published 27 June 1995. Four other applications relate to the spiro compounds WO 97/17350 published 15 May 1997; WO 97/34900 published 25 September 1997; WO 97/34901 published 25 September 1997; WO 97/35862

published 2 October 1997.

Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (c) are prepared according to the methods of international application publication number WO 95/30675, published 16 November 1995.

Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (f) are prepared according to the methods of international application publication number WO 95/17401, published 29 June 1995.

Suitably substituted anilines used to prepare compounds of formula (I) where E is a group or formula (g) are prepared according to the methods of international application publication number WO 96/31508 published 10 October 1996.

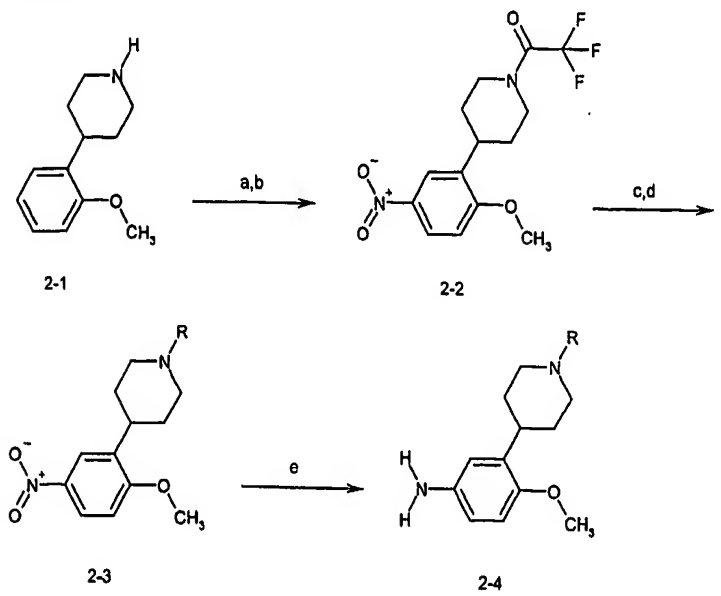
Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (h) are prepared according to the methods of international application publication number WO 95/32967, published 7 December 1995 and WO 97/07120, published 27 February 1997, WO 97/07120, published 27 February 1997.

Suitably substituted anilines used to prepare compounds of formula (I) where E is a group of formula (i) are prepared according to the methods of international application publication number WO 97/19070 published 29 May 1997.

Novel intermediates useful in preparing compounds of formula (I) are also included in the scope of this invention. For example, as shown in Scheme 2, certain anilines wherein E is a group (g) are prepared from commercially available 4-(2-methoxyphenyl)piperidine, **2-1** by treatment with a suitable acylating agent, for example trifluoroacetic anhydride, and suitable base, for example triethylamine, in a suitable solvent, for example dichloromethane. Nitration of the resulting N-acylated phenylpiperidine with a suitable nitrating agent, for example 70% nitric acid in acetic anhydride, at a suitable temperature, for example 0°C, for a suitable time, for example 30 minutes, yields **2-2**. Removal of the piperidine nitrogen protecting group from **2-2** with a suitable reagent, for example potassium carbonate, in a suitable solvent, for example aqueous methanol, at a suitable temperature, for example room temperature, gives **2-3** where R is H. Treatment of **2-3** where R is H with a suitable alkylating agent RX where R is C_{1-6}alkyl or C_{3-7}cycloalkyl, for example isopropyl, and X is a suitable leaving group, for example iodo, bromo, methanesulfonyloxy, trifluoromethysulfonyloxy, etc., and with a suitable base, for example potassium carbonate, in a suitable solvent, for example dimethylformamide and acetonitrile, at a suitable temperature, for example 70°C, for a suitable time, for example 20 hours gives **2-3** where R is C_{1-6}alkyl or C_{3-7}cycloalkyl. Alternatively, **2-3** where R is H may be reductively alkylated on the piperidine nitrogen by treatment with a C_{1-6}aldehyde, C_{3-6}ketone, or a C_{3-7}cyclic ketone, for example, cyclopentanone, and a suitable reducing

agent, for example sodium cyanoborohydride, in a suitable solvent, for example, acetic acid and methanol, for a suitable time, for example 16 hours, to afford **2-3** where R is C_{1-6}alkyl or C_{3-7}cycloalkyl. Reduction of the nitro group in **2-3** where R is C_{1-6}alkyl or C_{3-7}cycloalkyl with a suitable reagent, for example hydrogen, in the presence of a suitable catalyst, for example palladium hydroxide, in a suitable solvent, for example ethanol, for a suitable time, for example 4 hours, affords **2-4**. Compounds **2-4** are examples of **1-1** in Scheme 1 and are converted to **1-3**, which are compounds of formula (I)

Scheme 2



2-1 2-2 2-3 2-4

(a) TFAA, Et_3N, CH_2Cl_2, 16 h; (b) HNO_3, Ac_2O, 0°C, 30 min; (c) K_2CO_3, MeOH, H_2O, 40 h; (d) K_2CO_3, RX, DMF, MeCN, 70°C, 20 h or RCHO/RRCO, $NaBH_3CN$, AcOH, MeOH, Δ, 16 h; (e) H_2, $Pd(OH)_2$, EtOH, 4 h.

Particularly useful intermediates for preparing compounds of formula (I) are:
4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]benzenamine;
4-methoxy-3-[1-cyclopentyl-4-piperidinyl]benzenamine; and
4-methoxy-3-[1-(3-pentyl)-4-piperidinyl]benzenamine.

The invention will now be described by reference to the following examples which are merely illustrative and are not to be construed as a limitation of the scope of the present invention. In the Examples, mass spectra were performed upon a VG Zab mass spectrometer using fast atom bombardment, unless otherwise indicated.

EXAMPLES

Preparation 1

Preparation of 4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl)benzenamine

a) 4-(2-methoxyphenyl)-1-(trifluoroacetyl)piperidine

Trifluoroacetic anhydride (8.1 g, 39 mmol) was added portionwise over 10 min to a solution of commercially available 4-(2-methoxyphenyl)piperidine (6.7 g, 35 mmol), triethylamine (7.8 g, 77 mmol), and dichloromethane (100 mL) at RT. The reaction was maintained at RT for 16 h. The resultant mixture was washed with saturated sodium bicarbonate, saturated ammonium chloride, and with brine, dried ($MgSO_4$), and concentrated *in vacuo* to afford 10 g (99%) of the title compound as an amber oil. MS(ES) m/e 288.1 $[M+H]^+$.

b) 4-(2-methoxy-5-nitrophenyl)-1-(trifluoroacetyl)piperidine

Nitric acid (70%, 3.1 mL) was added portionwise to a solution of the compound of Preparation 1(a) (5.0 g, 17 mmol) in acetic anhydride (17 mL) at 0°C. The mixture was maintained at 0°C for an additional 30 min, combined with an identical concurrently run reaction, and poured into water (600 mL). The pH of the resultant mixture was adjusted to >9 by the addition of aqueous sodium carbonate followed by 10% sodium hydroxide. The resulting mixture was extracted with dichloromethane (2 × 400 mL) and the combined organic layers were washed with brine, dried ($MgSO_4$), and concentrated *in vacuo* to give 12 g (>100%) of a 2.2:1 mixture of the title compound and its 3-nitro isomer. The crude product was recrystallized from methanol (30 mL) to give 5.9 g (54%) of the title compound as off-white crystals. MS(ES) m/e 333.1 $[M+H]^+$.

c) 4-(2-methoxy-5-nitrophenyl)piperidine

Potassium carbonate (10 g, 74 mmol) was added to a solution of the compound of Preparation 1(b) (4.9 g, 15 mmol), methanol (100 mL) and water (7.5 mL). The resultant mixture was stirred at RT for 40 h, concentrated *in vacuo*, and the residue partitioned between water and dichloromethane. The layers were separated and aqueous layer was extracted with dichloromethane. The combined organic layers were washed with brine, dried ($MgSO_4$), and concentrated *in vacuo* to give 3.7 g (>100%) of the title compound as an off-white solid. MS(ES) m/e 237.2 $[M+H]^+$.

d) 4-(2-methoxy-5-nitrophenyl)-1-(1-methylethyl)piperidine

Potassium carbonate (8.6 g, 62 mmol) and isopropyl iodide (8.0 g, 47 mmol) were added to a solution of the compound of Preparation 1(c) (3.7 g, 16 mmol), dimethylformamide (10 mL) and acetonitrile (50 mL). The resultant mixture was heated at 70°C for 20 h, concentrated *in vacuo*, and the residue partitioned between water and dichloromethane. The aqueous phase was extracted with dichloromethane and the combined organic layers were washed with water (3 × 100 mL) and with brine, dried ($MgSO_4$), and concentrated *in vacuo* to provide 4.0 g (90%) of the title compound as a yellow solid. MS(ES) m/e 279.2 $[M+H]^+$.

e) 4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl)benzenamine

Palladium hydroxide on carbon (1.2 g, 20% dry weight) was added to a solution of the compound of Preparation 1(d) (4.0 g, 14 mmol) in ethanol (100 mL). The mixture was hydrogenated at 50 psi for 4 h, filtered through Celite®, and concentrated *in vacuo*. The residue was dissolved in ether (200 mL) and washed with 10% sodium carbonate and with water (2 × 100 mL). The ether solution was dried ($MgSO_4$) and concentrated *in vacuo* to provide 3.0 g (84%) of the title compound as a tan solid. MS(ES) m/e 249.2 $[M+H]^+$.

Preparation 2

Preparation of 1-(5,6,7,8-Tetrahydro-1-naphthalenyl)piperazine

Following the general procedure of Kuipers, et. al., J. Med. Chem., **1995**, *38*, 1942-1954, bis(chloroethyl)amine hydrochloride (2 g, 11.2 mmol) was added to a solution of 5,6,7,8-tetrahydro-1-naphthylamine (1.65 g, 11.2 mmol) in chlorobenzene (15 mL) and the mixture was heated to 135°C for 2 days. The mixture was cooled, concentrated *in vacuo*, and the residue was purified by flash chromatography (silica gel, 5% methanol/dichloromethane) to give the title compound as a tan solid which was further purified by HPLC (YMC CombiPrep ODS-A, 50 × 20 mm, 20 mL/min, A:0.1% trifluoroacetic acid in acetonitrile B:0.1% aqueous trifluoroacetic acid, A:10 to 90% during 10 min, UV detection at 254 nm) to give the title compound as a tan solid (0.25 g).

Preparation 3

Preparation of Ethyl 3-(1-piperazinyl)benzoate

Following the general procedure of Kato et. al., WO 9802432 and of Preparation 2, except substituting ethyl 3-aminobenzoate for 5,6,7,8-tetrahydro-1-naphthylamine, gave the title compound. MS(ES) m/e 235.2 $[M+H]^+$.

Preparation 4

Preparation of 4-Methoxy-3-[1-cyclopentyl-4-piperidinyl]benzenamine

a) 4-(2-methoxy-5-nitrophenyl)-1-(cyclopentyl)piperidine

A solution of the compound of Preparation 1(c) (3.4 g, 14.4 mmol) in methanol (21 mL) was treated with acetic acid (8.5 g, 0.14 mol), cyclopentanone (6.12 g, 71.4 mmol) and sodium cyanoborohydride (3.74 g, 57.8 mmol). The resulting mixture was heated to reflux for 16 h, concentrated *in vacuo*, and the residue was partitioned between dichloromethane and 2N sodium hydroxide. The organic phase was dried ($MgSO_4$) and concentrated *in vacuo* to afford the title compound.

b) 4-methoxy-3-[1-cyclopentyl-4-piperidinyl]benzenamine

Following the general procedure of Preparation 1(e), except substituting the compound of Preparation 4(a) for the compound of Preparation 1(d), gave the title compound.

Preparation 5

Preparation of 4-Methoxy-3-[1-(3-pentyl)-4-piperidinyl]benzenamine

The title compound is prepared following the procedure of Preparation 4(a)-4(b), except substituting 3-pentanone for cyclopentanone.

Example 1

Preparation of N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenyl-1,2,3,6-tetrahydropyridine-1-carboxamide

A solution of triphosgene (0.23 g, 0.77 mmol) in dichloromethane (25 mL) was stirred in an ice bath and treated with a solution of 4-phenyl-1,2,3,6-tetrahydropyridine hydrochloride (0.5 g, 2.6 mmol) and triethylamine (1 g, 10.2 mmol) in dichloromethane added dropwise. The ice bath was removed and the mixture was stirred for 30 min, treated with 3-(2-diisopropylamino)ethoxy-4-methoxyaniline (WO 95/15954)(0.68 g, 2.55 mmol), and stirred for 16 h. The mixture was diluted with dichloromethane (50 mL), extracted with 5% sodium carbonate, dried (Na_2SO_4), and concentrated *in vacuo*. The residue was chromatographed (silica gel, 8% methanol/dichloromethane saturated with ammonia) to give the title compound. MS(ES) m/e 452.0 $[M+H]^+$.

Example 2

Preparation of N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl) piperazine-1-carboxamide;

Triphosgene (74 mg, 0.25 mmol) was added to a solution of 3-(2-diisopropylamino)ethoxy-4-methoxyaniline (WO 95/15954)(200 mg, 0.75 mmol) and

dichloromethane (3 mL) and maintained at RT for 30 min. Triethylamine (0.30 g, 0.42 mL, 3.0 mmol) was added and the resulting mixture was stirred for 1 h, treated with 1-(2,3-dimethylphenyl)piperazine (0.11 g, 0.60 mmol), and the mixture stirred at RT for 16 h. The mixture was washed with water, dried ($MgSO_4$) and concentrated *in vacuo*. The crude product was purified by chromatography (silica gel, 20:1:0.04 dichloromethane:methanol:triethylamine) to give 205 mg (70%) of the title compound as an off-white powder. MS(ES) m/e 483.1 $[M+H]^+$.

Examples 3-22

Following the procedure of Example 2, except substituting 1-phenylpiperazine, 1-(2-methylphenyl)piperazine, 1-[2-(acetamidomethyl)phenyl]-piperazine(GB 2309458), 1-[3-(trifluoromethyl)phenyl]piperazine, 1-(2-methoxyphenyl)piperazine, 1-(2-chlorophenyl)piperazine, 1-(3-chlorophenyl)piperazine, 1-(4-chlorophenyl)piperazine, 1-(2,6-dimethylphenyl)piperazine, 1-(2,3-dichlorophenyl)piperazine, and 1-(3,4-dichlorophenyl)piperazine for 1-(2,3-dimethylphenyl)piperazine, gave the following compounds:

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenylpiperazine-1-carboxamide: MS(ES) m/e 454. 9 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 469.1 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-acetamido-methylphenyl) piperazine-1-carboxamide: MS(ES) m/e 525.9 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-trifluoromethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 522.8 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methoxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 485.0 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-chlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 488.9 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 488.8 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-chlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 488.8 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,6-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.1 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 522.9 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-

4-(3,4-dichlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 522.7 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-(3-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 499.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-phenylpiperazine-1-carboxamide: MS(ES) m/e 469.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dihydro-2(1H)-quinolinone-6-yl)piperazine-1-carboxamide: MS(ES) m/e 524.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-cyanophenyl)piperazine-1-carboxamide: MS(ES) m/e 480.2 $[M+H]^+$;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide: MS(ES) m/e 527.2 $[M+H]^+$; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide: MS(ES) m/e 527.2 $[M+H]^+$.

Example 23

Preparation of 1'-(1-Methylethyl)spiro[benzofuran-3(2H),4'-piperidin]-5-amine

a) 5- and 7-nitrospiro[benzofuran-3(2H),4'-piperidine]

A solution of 1'-methyl-5- and 7-nitrospiro[benzofuran-3(2H),4'-piperidine] (WO 96/11934) (3 g, 12 mmol) and diisopropylethylamine (2.5 g, 19 mmol) in 1,2-dichloroethane (80 mL) was treated with 1-chloroethyl chloroformate (2.3 g, 16 mmol) at RT, stirred for 1 h, and heated to reflux for 20 min. The mixture was cooled, concentrated *in vacuo*, and the residue was dissolved in methanol and heated to reflux for 2 h, concentrated *in vacuo*, and the residue was partitioned between dichloromethane (250 mL) and 5% sodium bicarbonate (50 mL). The organic phase was washed with 5% sodium bicarbonate (50 mL) and the combined aqueous phase was extracted with dichloromethane (2 X 50 mL). The combined organic phase was dried (Na_2SO_4) and concentrated to afford the title compound (2.65 g).

b) 1'-(tert-butoxycarbonyl)-5-nitrospiro[benzofuran-3(2H),4'-piperidine]

A solution of the compound of Preparation 2(a) (2.65 g, 1.13 mmol) in tetrahydrofuran (300 mL) was treated with di-tert-butyl dicarbonate (2.6 g, 12 mmol) and stirred at RT for 16 h. The mixture was concentrated *in vacuo* and the residue was

crystallized from methanol to afford the title compound (2.1 g).

c) 5-nitrospiro[benzofuran-3(2H),4'-piperidine]

A solution of the compound of Preparation 2(b)(2.1 g, 6.3 mmol) in dichloromethane (50 mL) and trifluoroacetic acid (10 mL) was kept at RT for 5 h, concentrated *in vacuo*, and the residue was partitioned between dichloromethane (300 mL) and 5% sodium bicarbonate. The organic phase was washed with 5% sodium bicarbonate and the combined aqueous washes were extracted with dichloromethane. The combined organic phase was dried (Na_2SO_4) and concentrated *in vacuo* to give the title compound (1.45 g). MS(ES) m/e 235.1 $[+H]^+$.

d) 1'-(1-methylethyl)-5-nitrospiro[benzofuran-3(2H),4'-piperidine]

A mixture of the compound of Preparation 2(c) (1.45 g, 6.2 mmol), powdered potassium carbonate (0.86 g, 6.2 mmol) and dimethylformamide (50 mL) containing 2-iodopropane (1.1 g, 6.4 mmol) was stirred and heated to 50°C for 4 h, treated with 2-iodopropane (0.17 g, 1 mmol) at 50°C for 90 min, and treated with 2-iodopropane (0.1 g, 1 mmol) at 50°C for 2 h. The mixture was concentrated *in vacuo* and the residue was partitioned between ethyl acetate (200 mL) and water (20 mL). The organic phase was washed, dried ($MgSO_4$), concentrated *in vacuo*, and the residue was chromatographed (silica gel, 5% methanol:dichloromethane) to give the title compound (0.85 g).

e) 1'-(1-methylethyl)spiro[benzofuran-3(2H),4'-piperidin]-5-amine

A solution of the compound of Preparation 2(d) (0.78 g, 2.8 mmol) in methanol (250 mL) containing 10% palladium-on-carbon (0.375 g) was shaken in a hydrogen atmosphere (40 psi) for 40 min, filtered, and concentrated *in vacuo* to afford the title compound (0.6 g).

Example 24

Following the procedure of Example 2, except substituting the compound of Example 23(e) for 3-(2-diisopropylamino)ethoxy-4-methoxyaniline, gave the following compound:

N-[2,3-dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 463.1 $[M+H]^+$.

Examples 25-46

Following the procedure of Example 2, except substituting 1-(3-methylphenyl)piperazine, 1-(4-methylphenyl)piperazine, 1-(2,5-dimethylphenyl)piperazine, 1-(3,4-dimethylphenyl)piperazine, 1-(3,5-dichlorophenyl)piperazine, 1-(3-methoxyphenyl)piperazine, 1-(3,5-

dimethoxyphenyl)piperazine, 1-[3-(ethoxycarbonyl)phenyl]piperazine, 1-(2-cyanophenyl)piperazine, 1-(4-cyanophenyl)piperazine, 1-(2-pyridinyl)piperazine, 1-(4-pyridinyl)piperazine, 1-[4-chloro-3-(trifluoromethyl)phenyl]piperazine, 1-[2-methyl-3-(trifluoromethyl)phenyl]piperazine, 1-(1-naphthalenyl)piperazine, 1-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine, 1-(1H-indol-4-yl)piperazine, 1-(4-methoxyphenyl)-3-methylpiperazine, 1-(5-chloro-2-methoxyphenyl)piperazine, 1-(3-hydroxyphenyl)piperazine, 1-(5-chloro-2-methylphenyl)piperazine, and 1-(3-chloro-2-methylphenyl)piperazine for 1-(2,3-dimethylphenyl)piperazine, gave the title compounds:

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 469.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 469.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,5-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethylphenyl)piperazine-1-carboxamide: MS(ES) m/e 483.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dichlorophenyl)piperazine-1-carboxamide: MS(ES) m/e 523.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methoxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 485.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethoxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 515.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide: MS(ES) m/e 527.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-cyanophenyl)piperazine-1-carboxamide: MS(ES) m/e 480.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-cyanophenyl)piperazine-1-carboxamide: MS(ES) m/e 480.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyridinyl)piperazine-1-carboxamide: MS(ES) m/e 456.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-pyridinyl)piperazine-1-carboxamide: MS(ES) m/e 456.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-chloro-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide: MS(ES) m/e 557.2 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide: MS(ES) m/e 537.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1-naphthalenyl)piperazine-1-carboxamide: MS(ES) m/e 505.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide: MS(ES) m/e 509.6 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1H-indol-4-yl)piperazine-1-carboxamide: MS(ES) m/e 494.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)-3-methylpiperazine-1-carboxamide: MS(ES) m/e 499.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 519.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-hydroxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 471.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 503.4 $[M+H]^+$; and

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide: MS(ES) m/e 503.4 $[M+H]^+$.

Example 47

Preparation of 4-(2,3-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide

Triphosgene (12.2 mg, 0.041 mmol) was added to a solution of the compound of Preparation 1(e) (31 mg, 0.125 mmol) in dichloromethane (1 mL). The mixture was stirred for 30 min and then triethylamine (0.07 mL, 0.5 mmol) was added. The mixture was stirred an additional 1 h, treated with 1-(2,3-dimethylphenyl)piperazine (31.0 mg, 0.125 mmol), and the mixture stirred at RT overnight. The resultant mixture was concentrated *in vacuo* and the residue was purified by preparative HPLC (YMC CombiPrep ODS-A, 50 × 20 mm, 20 mL/min, A:0.1% trifluoroacetic acid in acetonitrile B:0.1% aqueous trifluoroacetic acid, A:10 to 90% during 10 min, UV detection at 254 nm) to give 30 mg (52%) of the title compound as a yellow oil. MS(ES) m/e 465.4 $[M+H]^+$.

Example 48

Preparation of 4-(2,3-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide

Following the procedure of Example 47, except substituting 1-(2,3-dichlorophenyl)piperazine for 1-(2,3-dimethylphenyl)piperazine, gave the title compound. MS(ES) m/e 505.4 $[M+H]^+$.

Example 49

Preparation of N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-carboxyphenyl)piperazine-1-carboxamide

To a flask containing the compound of Example 32 (5.5 mg, 0.01 mmol) was added 0.5 ml ethanol and 0.3 N sodium hydroxide (0.1 ml, 0.03 mmol.). The mixture was stirred at RT overnight. The resultant mixture was concentrated *in vacuo* and the residue was purified by preparative HPLC (YMC CombiPrep ODS-A, 50 × 20 mm, 20 mL/min, A:0.1% trifluoroacetic acid in acetonitrile B:0.1% aqueous trifluoroacetic acid, A:10 to 90% during 10 min, UV detection at 254 nm) to give 1.0 mg (19%) of the title compound as a yellow oil. MS(ES) m/e 499.4 $[M+H]^+$.

Examples 50-51

Following the procedure of Example 49, except substituting the compounds of Examples 22 and 21 for the compound of Example 32 gave the title compounds:

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-carboxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 499.4 $[M+H]^+$; and

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-carboxyphenyl)piperazine-1-carboxamide: MS(ES) m/e 499.4 $[M+H]^+$.

Examples 52-61

Following the procedure of Example 2, except substituting 1-(3,4-dimethoxyphenyl)piperazine, 4-(2-benzothiazolyl)piperidine, 4-(1H-indol-2-yl)-1-piperidine, 4-(4-chlorophenyl)-4-hydroxy-1-piperidine, 4-acetyl-4-(4-chlorophenyl)-1-piperidine, 4-(4-chlorophenyl)-4-cyano-1-piperidine, 4-(4-hydroxyphenyl)-1-piperidine, 1-(6-chloro-2-benzothiazolyl)piperazine, 1-(2-pyrazinyl)piperazine, and 1-[5-(trifluoromethyl)-2-pyridinyl]piperazine for 1-(2,3-dimethylphenyl)piperazine, gave the following compounds:

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethoxyphenyl)-1-piperazinecarboxamide: MS(ES) m/e 515.4 $[M+H]^+$;

4-(2-benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide: MS(ES) m/e 511.4 $[M+H]^+$;

4-(1H-indol-2-yl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide: MS(ES) m/e 493.4 $[M+H]^+$;

4-(4-chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-4-hydroxy-1-piperidinecarboxamide: MS(ES) m/e 504.4 $[M+H]^+$;

4-acetyl-4-(4-chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-

methoxypheny]-1-piperidinecarboxamide: MS(ES) m/e 496.4 $[M+H]^+$;

4-(4-chlorophenyl)-4-cyano-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide: MS(ES) m/e 479.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-4-(4-hydroxyphenyl)--1-piperidinecarboxamide: MS(ES) m/e 470.4 $[M+H]^+$;

4-(6-chloro-2-benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperazinecarboxamide: MS(ES) m/e 546.4 $[M+H]^+$;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide: MS(ES) m/e 457.4 $[M+H]^+$; and

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide: MS(ES) m/e 524.4 $[M+H]^+$.

Examples 62-96

Following the procedure of Example 47, except substituting 4-(2-benzothiazolyl)-1-piperidine, 4-(1H-indol-2-yl)-1-piperidine, 4-(4-chlorophenyl)-4-hydroxy-1-piperidine, 4-acetyl-4-(4-chlorophenyl)-1-piperidine, 4-(4-chlorophenyl)-4-cyano-1-piperidine, 4-(4-hydroxyphenyl)-1-piperidine, 1-(6-chloro-2-benzothiazolyl)piperazine, 1-(2-pyrazinyl)piperazine, 1-[5-(trifluoromethyl)-2-pyridinyl]piperazine, 1-(3,4-dimethoxyphenyl)piperazine, 1-(2-chlorophenyl)piperazine, 1-(3-chlorophenyl)-piperazine, 1-(4-chlorophenyl)piperazine, 1-(3,4-dichlorophenyl)piperazine, 1-(3,5-dichlorophenyl)piperazine, 1-(2,6-dimethylphenyl)piperazine, 1-(2,4-dimethylphenyl)-piperazine, 1-(3,5-dimethylphenyl)piperazine, 1-(3-methylphenyl)piperazine, 1-(2,5-dimethylphenyl)piperazine, 1-(3,4-dimethylphenyl)piperazine, 1-(5,6,7,8-tetrahydro-1-naphthalenyl)piperazine, 1-(2-methylphenyl)piperazine, 1-(5-chloro-2-methylphenyl-piperazine, 1-(3-chloro-2-methylphenyl)piperazine, 1-(3-chloro-2-methoxyphenyl)-piperazine, 1-[3-(trifluoromethyl)phenyl]piperazine, 1-[4-chloro-3-(trifluoromethyl)-phenyl]piperazine, 1-[2-methyl-3-(trifluoromethyl)phenyl]piperazine, 1-(3-methoxyphenyl)piperazine, 1-(3,5-dimethoxyphenyl)piperazine, 1-(2-cyanophenyl)piperazine 1-(4-cyanophenyl)piperazine, the compound of Preparation 3, and 1-(1H-indol-4-yl)piperazine for 1-(2,3-dimethylphenyl)piperazine, gave the following compounds:

4-(2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide: MS(ES) m/e 493.4 $[M+H]^+$;

4-(1H-indol-2-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide: MS(ES) m/e 475.4 $[M+H]^+$;

4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-hydroxy-1-piperidinecarboxamide: MS(ES) m/e 486.4 $[M+H]^+$;

4-acetyl-4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-

piperidinyl]phenyl]-1-piperidinecarboxamide: MS(ES) m/e 478.4 $[M+H]^+$;

4-(4-chlorophenyl)-4-cyano-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide: MS(ES) m/e 461.4 $[M+H]^+$;

4-(4-hydroxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide: MS(ES) m/e 452.2 $[M+H]^+$;

4-(6-chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 528.2 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide: MS(ES) m/e 439.2 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide: MS(ES) m/e 506.4 $[M+H]^+$;

4-(3,4-dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 497.4 $[M+H]^+$;

4-(2-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 471.4 $[M+H]^+$;

4-(3-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 471.4 $[M+H]^+$;

4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 471.4 $[M+H]^+$;

4-(3,4-dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 505.4 $[M+H]^+$;

4-(3,5-dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 505.4 $[M+H]^+$;

4-(2,6-dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 465.4 $[M+H]^+$;

4-(2,4-dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 465.4 $[M+H]^+$;

4-(3,5-dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 465.4 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide: MS(ES) m/e 451.4 $[M+H]^+$;

4-(2,5-dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 465.4 $[M+H]^+$;

4-(3,4-dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 465.4 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide: MS(ES) m/e 491.4 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide: MS(ES) m/e 451.4 $[M+H]^+$;

4-(5-chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 485.4 $[M+H]^+$;

4-(3-chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 485.4 $[M+H]^+$;

4-(3-chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 501.4 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide: MS(ES) m/e 505.4 $[M+H]^+$;

4-[4-chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 539.4 $[M+H]^+$;

N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide: MS(ES) m/e 519.4 $[M+H]^+$;

4-(3-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 467.4 $[M+H]^+$;

4-(3,5-dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 497.4 $[M+H]^+$;

4-(2-cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 462.4 $[M+H]^+$;

4-(4-cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 462.4 $[M+H]^+$;

4-[3-(ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 509.4 $[M+H]^+$; and

4-(1H-indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide: MS(ES) m/e 476.4 $[M+H]^+$.

Example 97

Preparation of 4-[3-(Ethoxycarbonyl)phenyl]-N-[3-[3-[bis(1-methylethyl)amino]propoxy]-4-methoxyphenyl]-1-piperazinecarboxamide

Following the procedure of Example 2, except substituting 3-(3-diisopropylamino)propoxy-4-methoxyaniline (WO 99/01127) for 3-(2-diisopropylamino)ethoxy-4-methoxyaniline and substituting the compound of Preparation 3 for 1-(2,3-dimethylphenyl)piperazine, gave the title compound. MS(ES) m/e 541.4 $[M+H]^+$.

Example 98-99

Preparation of 4-(2,3-Dimethylphenyl)-N-[4-methoxy-3-[1-cyclopentyl-4-piperidinyl]phenyl]-1-piperazinecarboxamide and 4-(2,3-Dimethylphenyl)-N-[1-(3-pentyl)-4-methoxy-3-[1-cyclopentyl-4-piperidinyl]phenyl]-1-piperazinecarboxamide

Following the general procedure of Example 47, except substituting the compounds of Preparation 4(b) and Preparation 5 for the compound of Preparation 1(e), gives the title compounds.

Biological Data:

CCR5 Receptor Binding Assay

CHO cell membranes (0.25×10^6 cell equivalents) derived from CHO cells stably transfected with CCR5 were incubated with 0.3 ^{125}I-RANTES in a 96 well plate for 45 min. at room temperature (final reaction volume 200 uL). The reaction was terminated by filtration and the filters (GF/C) were washed twelve times with a solution of phosphate buffered saline containing 0.1 % bovine serum albumin and 0.05 % NaN_3. The radioactivity bound to filters was measured by liquid scintillation spectrometry. Non-specific binding was determined in the presence of unlabelled RANTES (10 or 30 nM) and averages 30-50% of total binding.

CCR5 Receptor Functional Assay

The cellular functional assay used to assess antagonist activity of compounds was RANTES-induced Ca^{2+} mobilization in RBL 2H3 cells stably expressing the hCCR5 receptor (RBL 2H3 hCCR5). Agonist activity is determined by Ca^{2+} mobilization in the same cells which is inhibitable by a selective CCR5 antagonist. Cells were grown to 80-100% confluency in T-150 flasks and washed with phosphate-buffered saline. Cells were lifted from the flasks by treating with 3 mL of 1 mM EDTA for 3 min. at room temperature and diluting to 2×10^6 cells/mL with Krebs Ringer Henseleit buffer (KRH; 118 mM NaCl, 4.6 mM KCl, 25 mM $NaHCO_3$, 1 mM KH_2PO_4 and 11 mM glucose) containing 5 mM HEPES (pH 7.4), 1 mM $CaCl_2$, 1 mM $MgCl_2$ and 0.1% BSA and centrifuged at 200g for 3 min. Cells were resuspended at 2×10^6 cells/mL in the same buffer with 2 µM Fura-2AM, and incubated for 35 min. at 37° C. Cells were centrifuged at 200 x g for 3 min. and resuspended in the same buffer without Fura-2AM, then incubated for 15 min. at 37° C to complete the hydrolysis of intracellular Fura-2AM, and then centrifuged as before. Cells (10^6 cells/mL) were resuspended in cold KRH with 5 mM HEPES (pH 7.4), 1 mM $CaCl_2$, 1 mM $MgCl_2$ and 0.1% gelatin and maintained on ice until assayed. For antagonist studies, aliquots (2 mL) of cells were pre-warmed at 37° C for 5 min. in 3 mL plastic cuvettes and fluorescence

measured in a fluorometer (Johnson Foundation Biomedical Group, Philadelphia, PA, USA) with magnetic stirring and temperature maintained at 37° C. Excitation was set at 340 nm and emission set at 510 nm. Various concentrations of antagonists or vehicle were added and fluorescence monitored for ~15 sec to ensure that there was no change in baseline fluorescence, followed by the addition of 33 nM RANTES. Maximal Ca^{2+} attained after 33 nM RANTES stimulation was calculated as described by Grynkiewicz *et al.*, (1985). The percent of maximal RANTES-induced Ca^{2+} was determined for each concentration of antagonist and the IC_{50}, defined as the concentration of test compound that inhibits 50% of the maximal 33 nM RANTES response, obtained from the concentration-response curves (5-7 concentrations of antagonists).

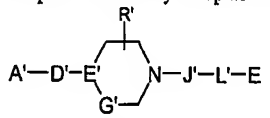
The compounds of this invention show CCR5 receptor modulator activity having IC_{50} values in the range of 0.0001 to 100 μM. The full structure/activity relationship has not yet been established for the compounds of this invention. However, given the disclosure herein, one of ordinary skill in the art can utilize the present assays in order to determine which compounds of formula (I) are modulators of the CCR5 receptor and which bind thereto with an IC_{50} value in the range of 0.0001 to 100 μM.

All publications, including, but not limited to, patents and patent applications cited in this specification, are herein incorporated by reference as if each individual publication were specifically and individually indicated to be incorporated by reference herein as though fully set forth.

The above description fully discloses the invention including preferred embodiments thereof. Modifications and improvements of the embodiments specifically disclosed herein are within the scope of the following claims. Without further elaboration it is believed that one skilled in the art can, given the preceding description, utilize the present invention to its fullest extent. Therefore any examples are to be construed as merely illustrative and not a limitation on the scope of the present invention in any way. The embodiments of the invention in which an exclusive property or privilege is claimed are defined as follows.

What is claimed is:

1. A method of treating a CCR5-mediated disease state in mammals which comprises administering to a mammal in need of such treatment, an effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof:



Formula I

in which:

the basic nitrogen in moiety E may be optionally quaternized with C_{1-6}alkyl or is optionally present as the N-oxide;

A' is aryl or heteroaryl, each of which is optionally substituted with one or more of $R^{1'}$; or A' is aryl or heteroaryl fused to a saturated or partly unsaturated 5-7-membered ring to form a higher order ring moiety, which ring moiety optionally contains 1 or 2 heteroatoms selected from oxygen, nitrogen or sulfur, wherein nitrogen may be optionally substituted with hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl; wherein the higher order ring moiety is optionally substituted with one or more of $R^{1'}$;

$R^{1'}$ is hydrogen, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, CH_2CF_3, aryl, aralkyl, $(CH_2)_{a'}NR^{2'}R^{3'}$, $(CH_2)_{a'}NR^{2'}COR^{4'}$, $(CH_2)_{a'}NR^{2'}CO_2R^{5'}$, $(CH_2)_{a'}NR^{2'}SO_2R^{6'}$, $(CH_2)_{a'}CONR^{7'}R^{8'}$, hydroxy$C_{1-6}$alkyl, C_{1-4}alkoxyalkyl (optionally substituted by a C_{1-4}alkoxy or hydroxy group), $(CH_2)_{a'}CO_2C_{1-6}$alkyl, $(CH_2)_{b'}OC(O)R^{9'}$, $CR^{10'}{=}NOR^{11'}$, $CNR^{10'}{=}NOR^{11'}$, $COR^{12'}$, $CONR^{7'}R^{8'}$, $CONR^{7'}(CH_2)_{c'}OC_{1-4}$alkyl, $CONR^{7'}(CH_2)_{a'}CO_2R^{13'}$, $CONHNR^{14'}R^{15'}$, $CONR^{7'}SO_2R^{16'}$, $CO_2R^{17'}$, cyano, trifluoromethyl, $NR^{2'}R^{3'}$, $NR^{2'}COR^{4'}$, $NR^{18'}CO(CH_2)_{a'}NR^{18'}R^{19'}$, $NR^{18'}CONR^{18'}R^{19'}$, $NR^{2'}CO_2R^{5'}$, $NR^{2'}SO_2R^{6'}$, $N{=}CNR^{18'}NR^{18'}R^{19'}$, nitro, hydroxy, C_{1-6}alkoxy, OCF_3, hydroxyC_{1-6}alkoxy, C_{1-6}alkoxyC_{1-6}alkoxy, $OC(O)NR^{20'}R^{21'}$, $SR^{22'}$, $SOR^{23'}$, $SO_2R^{23'}$, $SO_2NR^{20'}R^{21'}$ or halogen, or $R^{1'}$ is a 5- to 7-membered ring containing 1 to 4 heteroatoms selected from nitrogen, oxygen, or sulfur, optionally substituted with hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, C_{3-6}cycloalkenyl, hydroxyC_{1-6}alkyl, (C_{1-6}alkyl)C_{1-6}alkyl, $CONR^{7'}R^{8'}$, $CO_2R^{17'}$, cyano, aryl, trifluoromethyl, nitro, hydroxy, C_{1-6}alkoxy, acyloxy, or halogen;

a' is 1, 2, 3 or 4;

b' is 0, 1, 2 or 3;

c' is 1, 2 or 3;

$R^{2'}$ and $R^{3'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{2'}$ and $R^{3'}$ together with the nitrogen to which they are attached, form a 5- to 6-membered heterocyclic ring which ring may be optionally substituted by an oxo group, or, when there are 6 ring members, the ring may optionally contain one oxygen or one sulfur atom;

$R^{4'}$ is hydrogen, C_{1-6}alkyl or C_{1-4}alkoxyalkyl, or, when $R^{1'}$ is $NR^{2'}COR^{4'}$, $R^{4'}$ is $(CH_2)_{1-3}$ and forms a ring with A';

$R^{5'}$ is C_{1-6}alkyl;

$R^{6'}$ is C_{1-6}alkyl or phenyl;

$R^{7'}$ and $R^{8'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{7'}$ and $R^{8'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring, wherein when there are 6 ring members, the ring may optionally contain one oxygen or one sulfur atom;

$R^{9'}$ is C_{1-4}alkyl, optionally substituted by a C_{1-6}alkoxy;

$R^{10'}$ and $R^{11'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{12'}$ is hydrogen or C_{1-6}alkyl;

$R^{13'}$ is hydrogen or C_{1-6}alkyl;

$R^{14'}$ and $R^{15'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{16'}$ is hydrogen or C_{1-6}alkyl;

$R^{17'}$ is hydrogen or C_{1-6}alkyl optionally substituted with one or more substituents selected from C_{1-6}alkyl, C_{1-6}alkoxy, hydroxy, or $NR^{2'}R^{3'}$;

$R^{18'}$ and $R^{19'}$ are independently hydrogen or C_{1-6}alkyl;

$R^{20'}$ and $R^{21'}$ are independently hydrogen or C_{1-6}alkyl, or $R^{20'}$ and $R^{21'}$ together with the nitrogen to which they are attached form a 5- to 6-membered saturated heterocyclic ring which, when the ring is 6-membered, may optionally contain in the ring one oxygen or one sulfur atom.

$R^{22'}$ is hydrogen or C_{1-6}alkyl;

$R^{23'}$ is C_{1-6}alkyl;

D' is either a bond or represents $[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{a''}CO$, CO, SO_2, $CO[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}$, $S[C(R^{24'})_2]_{a''}$, $O[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}OCO$, $NR^{25'}[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}CO$, $[C(R^{24'})_2]_{c''}NR^{25'}CO$, $NR^{25''}CO[C(R^{24'})_2]_{a''}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}$, $[C(R^{24'})_2]_{c''}NR^{25'}SO_2$, $CR^{24'}=CR^{24'}CO$, C≡CCO, $(C(R^{24'})_2)_{c''}SO_2$, $SO_2[C(R^{24'})_2]_{a''}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}SO_2$, $O[C(R^{24'})_2]_{a''}SO_2$, $SO_2NR^{25'}[C(R^{24'})_2]_{1-2}$, $[C(R^{24'})_2]_{b''}COO[C(R^{24'})_2]_2$, $[C(R^{24'})_2]_{b''}CONR^{25'}[C(R^{24'})_2]_{1-2}$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$, then D' may further be O, $NR^{25'}$, $CONR^{25'}$, $SO_2NR^{25'}$, $OCONR^{25'}$, $NR^{25'}COO$, $NR^{25'}CONR^{25'}$, $[C(R^{24'})_2]_{a''}NR^{25'}[C(R^{24'})_2]_{b''}$, $[C(R^{24'})_2]_{a''}O[C(R^{24'})_2]_{b''}$,

$CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}O$, $NR^{25'}[C(R^{24'})_2]_{a''}NR^{25'}$, $O[C(R^{24'})_2)]_{a''}NR^{25'}$, $O[C(R^{24'})_2]_{a''}O$, $CO[C(R^{24'})_2]_{a''}O$, $SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $SO_2[C(R^{24'})_2]_{a''}O$, $[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $[C(R^{24'})_2]_{a''}CONR^{25'}$, $O[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $O[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}SO_2NR^{25'}$, $NR^{25'}[C(R^{24'})_2]_{a''}CONR^{25'}$, $NR^{25'}CO[C(R^{24'})_2]_{a''}NR^{25'}$, $NR^{25'}SO_2[C(R^{24'})_2]_{a''}NR^{25'}$, $(C(R^{24'})_2)_{a''}S(C(R^{24'})_2)_{b''}$, COO, $CR^{24'}OH$, $C(R^{24'})_{a''}CR^{24'}OH$; and when E' and G' together are $CR^{27'}$-$C(R^{26'})_2$ or $C=CR^{26'}$, then D' may further be $CR^{24'}=CR^{24'}$ or C≡C; wherein a'' is 1-6, b'' is 0-1, and c'' is 0-2;

$R^{24'}$ is hydrogen or C_{1-6}alkyl;

$R^{25'}$ is hydrogen or C_{1-6}alkyl;

E' and G' together are $NC(R^{26'})_2$, $NC(R^{26'})_2C(R^{26'})_2$, $CR^{27'}C(R^{26'})_2$ or $C=CR^{26'}$;

$R^{26'}$ is hydrogen or C_{1-6}alkyl;

$R^{27'}$ is hydrogen, $OR^{28'}$, $NHR^{28'}$, CN, NO_2, $R^{28'}$, $SR^{29'}$, $COR^{28'}$, $CHOHR^{28'}$, $CO_2R^{28'}$, $NHCOR^{28'}$, $NHCO_2R^{29'}$, $NHSO_2R^{29'}$, or $OCONHR^{28'}$;

$R^{28'}$ is hydrogen, C_{1-5}alkyl, aryl or aralkyl;

$R^{29'}$ is C_{1-5}alkyl, aryl or aralkyl;

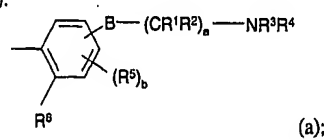
R' is one or more of hydrogen or C_{1-6}alkyl, or R' is oxo;

J' is CO or SO_2;

L' is $NR^{30'}$, O or $C(R^{30'})_2$;

$R^{30'}$ is hydrogen or C_{1-6}alkyl;

E represents a group (a):



(a);

in which

B is oxygen, C≡C, $S(O)_c$, $CR^7=CR^8$, or CR^7R^8, or B is NR^9;

R^1 and R^2 are independently hydrogen or C_{1-6}alkyl; alternatively $B(CR^1R^2)_a$ is $OCR^1R^2CR^1(OH)CR^1R^2$ or $OCR^1R^2CR^1(OCOCH_3)CR^1R^2$;

R^3 and R^4 are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{10}R^{11}$, $NR^{10}R^{11}$, hydroxy, $OCOR^{12}$, $NHCOCF_3$, $NHSO_2R^{13}$, $NHCO_2R^{14}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is

optionally substituted by OH;

R^5 is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{15}R^{16}$, CO_2R^{17}, trifluoromethyl, $NHCO_2R^{18}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_dR^{19}$, $SO_2NR^{20}R^{21}$ or halogen;

R^6 is hydrogen, C_{1-6}alkyl, aryl, trifluoromethyl, hydroxy, C_{1-6}alkoxy or halogen, or R^6 taken together with $R^{30'}$ forms a group D where D is $(CR^{22}R^{23})_e$ or D is $(CR^{22}R^{23})_f$-G where G is oxygen, sulfur or $CR^{22}=CR^{23}$, $CR^{22}=N$, $=CR^{22}O$, $=CR^{22}S$, or $=CR^{22}-NR^{23}$;

R^7, R^8, R^{10}, R^{11}, R^{12}, R^{15}, R^{16}, R^{17}, R^{20}, R^{21}, R^{22}, and R^{23} are independently hydrogen or C_{1-6}alkyl;

R^9 is hydrogen, C_{1-6}alkyl, or phenylC_{1-6}alkyl;

R^{13}, R^{14}, R^{18}, and R^{19} are independently C_{1-6}alkyl;

a is 1, 2, 3, or 4;

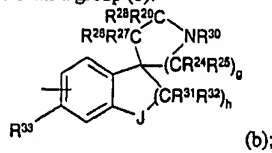
b is 1 or 2;

c and d are independently 0, 1 or 2;

e is 2, 3 or 4;

f is 0, 1, 2 or 3;

alternatively, E represents a group (b):



(b);

R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{31}, and R^{32} are independently hydrogen or C_{1-6}alkyl;

R^{30} is hydrogen, C_{1-6}alkyl, or C_{3-7}cycloalkyl;

R^{33} is hydrogen, C_{1-6}alkyl, trifluoromethyl, hydroxy or halogen, or R^{33} and $R^{30'}$ together form a group -K- where K is $(CR^{34}R^{35})_i$ or K is $(CR^{34}R^{35})_j$ -M and M is oxygen, sulfur, $CR^{34}=CR^{35}$, $CR^{34}=N$, or N=N;

J is oxygen, $CR^{36}R^{37}$, or NR^{38}, or J is a group $S(O)_k$;

R^{34}, R^{35}, R^{36}, R^{37}, and R^{38} are independently hydrogen or C_{1-6}alkyl;

g is 1, 2 or 3;

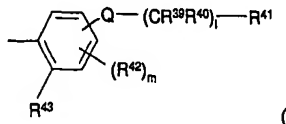
h is 1, 2 or 3;

i is 2, 3, or 4;

j is 0, 1, 2, or 3;

k is 0, 1 or 2;

alternatively, E represents a group (c):



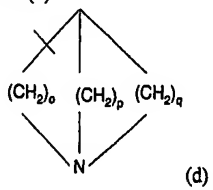
(c);

in which:

Q is oxygen, $S(O)_n$, $CR^{44}=CR^{45}$, $CR^{44}R^{45}$, or Q is NR^{46};

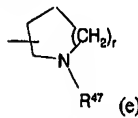
R^{39} and R^{40} are independently hydrogen or C_{1-6}alkyl;

R^{41} is a group of formula (d):



(d)

or R^{41} is a group of formula (e):



(e)

R^{42} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{48}R^{49}$, CO_2R^{50}, trifluoromethyl, $NHCO_2R^{51}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_sR^{52}$, $SO_2NR^{53}R^{54}$, or halogen;

R^{43} is hydrogen or R^{43} together with $R^{30'}$ forms a group R where R is $CR^{55}=CR^{56}$, $CR^{55}=CR^{56}CR^{55}R^{56}$, or $(CR^{55}R^{56})_t$;

R^{44}, R^{45}, R^{46}, R^{48}, R^{49}, R^{50}, R^{53}, R^{54}, R^{55}, and R^{56} are independently hydrogen or C_{1-6}alkyl;

R^{47} is hydrogen, C_{1-6}alkyl, or C_{3-7} cycloalkyl;

R^{51} and R^{52} are independently C_{1-6}alkyl;

l is 0, 1, 2, or 3;

m is 1 or 2;

n is 0, 1, or 2

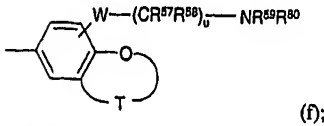
o, p, and q are independently integers having the value 1, 2, or 3;

r is 0,1, 2, or 3;

s is 0, 1, or 2;

t is 2 or 3;

alternatively, E represents a group (f):



(f);

R^{57} and R^{58} are independently hydrogen or C_{1-6}alkyl;

R^{59} and R^{60} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{61}R^{62}$, $NR^{61}R^{62}$, hydroxy, $OCOR^{63}$, $NHCOCF_3$, $NHSO_2R^{64}$, $NHCO_2R^{65}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

T is $-(CR^{66}R^{67})_v-$ or $-O(CR^{66}R^{67})_w-$;

W is oxygen, $S(O)_x$, NR^{68}, or W is $CR^{69}=CR^{70}$ or $CR^{69}R^{70}$;

R^{61}, R^{62}, R^{63}, R^{66}, R^{67} R^{68}, R^{69}, and R^{70} are independently hydrogen or C_{1-6}alkyl;

R^{64} and R^{65} are independently C_{1-6}alkyl;

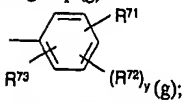
u is 1 to 4;

v is 2 or 3;

w is 1, 2, or 3;

x is 0, 1 or 2;

alternatively, E represents a group (g):



(g);

R^{71} is a 5- to 7-membered saturated or partially saturated heterocyclic ring containing a nitrogen atom and optionally a further 1 or 2 heteroatoms selected from nitrogen, oxygen or sulfur or R^{71} is an optionally substituted 6,6 or 6,5 bicyclic ring containing a nitrogen atom and optionally a further heteroatom selected from oxygen, nitrogen or sulfur, which ring systems may be optionally substituted with one or more of C_{1-6}alkyl and optionally substituted on nitrogen with hydrogen, C_{1-6}alkyl or C_{3-7}cycloalkyl;

R^{72} is hydrogen, C_{1-6}alkyl, aryl, CN, $CONR^{74}R^{75}$, CO_2R^{76}, trifluoromethyl, $NHCO_2R^{77}$, hydroxy, C_{1-6}alkoxy, benzyloxy, $OCH_2CO_2C_{1-6}$alkyl, OCF_3, $S(O)_zR^{78}$, $SO_2NR^{79}R^{80}$, or halogen;

R^{73} is hydrogen, C_{1-6}alkyl, hydroxy, C_{1-6}alkoxy or halogen, or R^{73} and $R^{30'}$ taken together from a group -X- where X is $(CR^{81}R^{82})_{aa}$ or X is $(CR^{81}R^{82})_{ab}$-Y and

Y is oxygen, sulfur or $CR^{81}=CR^{82}$;

R^{74}, R^{75}, R^{76}, R^{79}, R^{80}, R^{81}, and R^{82} are independently hydrogen or C_{1-6}alkyl;

R^{77} and R^{78} are independently C_{1-6}alkyl;

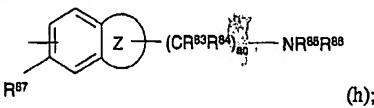
y is 1 or 2;

z is 0, 1, or 2;

aa is 2, 3 or 4;

ab is 0, 1, 2 or 3;

alternatively, E represents a group (h):



(h);

R^{83} and R^{84} are independently hydrogen or C_{1-6}alkyl;

R^{85} and R^{86} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{88}R^{89}$, $NR^{90}R^{91}$, hydroxy, $OCOR^{92}$, $NHCOCF_3$, $NHSO_2R^{93}$, $NHCO_2R^{94}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

R^{87} is hydrogen or C_{1-6}alkyl, C_{1-6}alkoxy, or halogen, or R^{87} together with $R^{30'}$ forms a group -AA- where AA is $(CR^{95}R^{96})_{ad}$ or AA is $(CR^{95}=CR^{96})_{ae}$-AB and AB is oxygen, sulfur, $CR^{95}=CR^{96}$, $CR^{95}=N$, $CR^{95}NR^{96}$ or N=N;

Z is an optionally substituted 5 to 7-membered heterocyclic ring containing 1 to 3 heteroatoms selected from oxygen, nitrogen or sulfur;

R^{88}, R^{89}, R^{90}, R^{91}, R^{92}, R^{95}, and R^{96} are independently hydrogen or C_{1-6}alkyl;

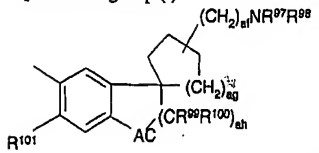
R^{93} and R^{94} are independently C_{1-6}alkyl;

ac is 0 to 4;

ad is 1, 2 or 3;

ae is 0, 1 or 2;

alternatively, E represents a group (i):



(i);

R^{97} and R^{98} are independently hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aralkyl, or together with the nitrogen atom to which they are attached form an optionally substituted 5- to 7-membered heterocyclic ring which may contain an additional heteroatom selected from oxygen, nitrogen or sulfur, where optional substituents include C_{1-6}alkyl, aryl, $CONR^{102}R^{103}$, $NR^{104}R^{105}$, hydroxy, $OCOR^{106}$, $NHCOCF_3$, $NHSO_2R^{107}$, $NHCO_2R^{108}$, or $NHCOC_{0-6}$alkyl wherein the alkyl of $NHCOC_{0-6}$alkyl is optionally substituted by OH;

R^{99} and R^{100} are independently hydrogen or C_{1-6}alkyl;

R^{101} is hydrogen or C_{1-6}alkyl or R^{101} and $R^{30'}$ together form a group -AD- where AD is $(CR^{109}R^{110})_{ai}$ or AD is $(CR^{109}R^{110})_{aj}$-AE and AE is oxygen, sulfur or $CR^{109}=CR^{110}$;

AC is oxygen, $CR^{111}R^{112}$ or NR^{113} or AC is a group $S(O)_{ak}$;

R^{102}, R^{103}, R^{104}, R^{105}, R^{106}, R^{109}, R^{110}, R^{111}, R^{112}, and R^{113} are independently hydrogen or C_{1-6}alkyl;

R^{107} and R^{108} are independently C_{1-6}alkyl;

af is 0, 1, 2, 3, or 4;

ag is 1, 2, or 3;

ah is 1, 2, 3 or 4;

ai is 2, 3 or 4;

aj is 0, 1, 2, or 3; and

ak is 0, 1 or 2.

2. The method as claimed in claim 1, wherein the compound of formula (I) is selected from:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenyl-1,2,3,6-tetrahydropyridine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-acetamido-methylphenyl) piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-

4-(3-trifluoromethylphenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,6-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dihydro-2(1H)-quinolinone-6-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[2,3-dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-chloro-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1-naphthalenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1H-indol-4-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)-3-methylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-hydroxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-

methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide;

4-(2,3-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,3-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethoxyphenyl)-1-piperazinecarboxamide;

4-(2-Benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(2-Benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-4-hydroxy-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-hydroxy-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxypheny]-4-(4-hydroxyphenyl)--1-piperidinecarboxamide;

4-(4-Hydroxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperazinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

4-(3,4-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,6-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-

tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-carboxyphenyl)piperazine-1-carboxamide.

3. The method as claimed in claim 1, wherein the compound of formula (I) is selected from:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-

dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[2,3-Dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[4-cyano-1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide.

4. The method as claimed in claim 1, wherein the compound of formula (I) is selected from:

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide; and

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide.

5. The method as claimed in claim 1, wherein the disease is selected from COPD, asthma and atopic disorders (for example, atopic dermatitis and allergies), rheumatoid arthritis, sarcoidosis, or idiopathic pulmonary fibrosis and other fibrotic diseases, atherosclerosis, psoriasis, autoimmune diseases such as multiple sclerosis, treating and/or preventing rejection of transplanted organs, inflammatory bowel disease, and HIV infection.

6. The method as claimed in claim 1, wherein A' is phenyl, 5,6,7,8-tetrahydro-1-naphthalenyl, or 1H-indol-4-yl; D' is a bond, E' and G' together are NCH_2, R' is hydrogen, J' is CO, L' is NH, and E is group (g).

7. The method as claimed in claim 6, wherein A' is phenyl, and $R^{1'}$ is methyl, chloro or trifluoromethyl substituted at the 2 and/or 3-positions, or $R^{1'}$ is 2,4-dimethyl, 2-methoxy-5-chloro, 2-methyl, 3-ethoxycarbonyl, or 3,5-dichloro.

8. A compound or a pharmaceutically active salt or solvate thereof, selected from:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenyl-1,2,3,6-tetrahydropyridine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-acetamido-

methylphenyl) piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-trifluoromethylphenyl)piperazine-1-carboxamide;

N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-chlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,6-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-3-methyl-4-phenylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dihydro-2(1H)-quinolinone-6-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[2,3-dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,5-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,5-dimethoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-cyanophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-pyridinyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[4-chloro-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1-naphthalenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(1H-indol-4-yl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-methoxyphenyl)-3-methylpiperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-

hydroxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide;

4-(2,3-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,3-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-carboxyphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3,4-dimethoxyphenyl)-1-piperazinecarboxamide;

4-(2-Benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(2-Benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperidinecarboxamide;

4-(1H-Indol-2-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-4-hydroxy-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-hydroxy-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide;

4-Acetyl-4-(4-chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[3-(2-diisopropylamino)ethoxy-4-methoxypheny]-1-piperidinecarboxamide;

4-(4-Chlorophenyl)-4-cyano-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxypheny]-4-(4-hydroxyphenyl)-1-piperidinecarboxamide;

4-(4-Hydroxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperidinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[3-(2-diisopropylamino)ethoxy-4-methoxyphenyl]-1-piperazinecarboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-pyrazinyl)-1-piperazinecarboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[5-(trifluoromethyl)-2-pyridinyl]-1-piperazinecarboxamide;

4-(3,4-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,6-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-

piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(4-carboxyphenyl)piperazine-1-carboxamide.

9. A compound or a pharmaceutically active salt or solvate thereof, selected from:

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-

dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,3-dichlorophenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(2,4-dimethylphenyl)piperazine-1-carboxamide;

N-[2,3-Dihydro-1'-isopropyl-spiro[benzofuran-5-yl-3,4'-piperidine]]-4-(2,3-dimethylphenyl)piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[3-(ethoxycarbonyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide;

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(5-chloro-2-methoxyphenyl)piperazine-1-carboxamide;

4-(6-Chloro-2-benzothiazolyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(4-Chlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(3-methylphenyl)-1-piperazinecarboxamide;

4-(2,5-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,4-Dimethylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(2-methylphenyl)-1-piperazinecarboxamide;

4-(5-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3-Chloro-2-methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[3-(trifluoromethyl)phenyl]1-piperazinecarboxamide;

4-[4-Chloro-3-(trifluoromethyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide;

4-(3-Methoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(3,5-Dimethoxyphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(2-Cyanophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[4-cyano-1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-[3-(Ethoxycarbonyl)phenyl]-N-[3-(3-diisopropylamino)propoxy-4-methoxyphenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-(3-chloro-2-methylphenyl)piperazine-1-carboxamide.

10. A compound or a pharmaceutically active salt or solvate thereof, selected from:

4-(3,5-Dichlorophenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-(5,6,7,8-tetrahydro-1-naphthalenyl)-1-piperazinecarboxamide;

4-(3-Chloro-2-methylphenyl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide;

N-[4-Methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-4-[2-methyl-3-(trifluoromethyl)phenyl]-1-piperazinecarboxamide; and

4-(1H-Indol-4-yl)-N-[4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]phenyl]-1-piperazinecarboxamide; and

N-[3-(2-Diisopropylamino)ethoxy-4-methoxyphenyl]-4-[1-(5,6,7,8-tetrahydronaphthalenyl]piperazine-1-carboxamide.

11. A pharmaceutical composition comprising a compound as claimed in claim 8, 9 or 10, and a pharmaceutically acceptable carrier.

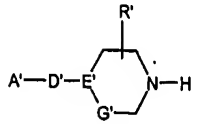
12. A process for making a compound as claimed in claims 8, 9, or 10, comprising for compounds wherein L' is $NR^{30'}$,

a) treating a compound of formula (II):

Formula (II)

wherein $R^{30'}$ is hydrogen or C_{1-6}alkyl, with triphosgene under basic conditions to form a mixture; and

b) adding to the mixture a compound of formula (III):



wherein A', D', E', G' and R' are as defined in claim 1.

13. A compound selected from:

4-methoxy-3-[1-(1-methylethyl)-4-piperidinyl]benzenamine;

4-methoxy-3-[1-cyclopentyl-4-piperidinyl]benzenamine; and

4-methoxy-3-[1-(3-pentyl)-4-piperidinyl]benzenamine.

INTERNATIONAL SEARCH REPORT

International application No. PCT/US01/22529

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) :A61K 31/495

US CL : 514/255

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/255

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Please See Extra Sheet.

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,789,412 A (HALAZY et al) 04 August 1998, see entire text.	1-10
A	WO 99/17773 A1(SMITHKLINE BEECHAM CORPORATION) 15 April 1999, see entire text.	1-10

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
18 SEPTEMBER 2001	30 OCT 2001
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer VICKIE KIM Telephone No. (703) 305-3667

Form PCT/ISA/210 (second sheet) (July 1998)*

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US01/22589

B. FIELDS SEARCHED
Electronic data bases consulted (Name of data base and where practicable terms used):

CAS ONLINE, REGISTRY, CAPLUS, USPATFUL,
search structure and terms:CCR5 1, arthritis, sarcoidosis, fibrosis, artherosclerosis, autoimmune disease, inflammatory bowel disease